

FORM PTO-1390 (Modified) (REV 11-98)		U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE		ATTORNEY'S DOCKET NUMBER 37921-151956
TRANSMITTAL LETTER TO THE UNITED STATES DESIGNATED/ELECTED OFFICE (DO/EO/US) CONCERNING A FILING UNDER 35 U.S.C. 371				U.S. APPLICATION NO (IF KNOWN, SEE 37 CFR 09/831686
INTERNATIONAL APPLICATION NO. PCT/AU99/00993	INTERNATIONAL FILING DATE November 10, 1999	PRIORITY DATE CLAIMED November 11, 1998		
TITLE OF INVENTION Biological Compositions, Components Thereof and Uses Therefor				
APPLICANT(S) FOR DO/EO/US Stephen Alister Locarnini, Joseph Torresi, Linda Ernest-Silveira and Ingrid Angeline Bartholmeusz				
Applicant herewith submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information:				
<ol style="list-style-type: none"> <input checked="" type="checkbox"/> This is a FIRST submission of items concerning a filing under 35 U.S.C. 371. <input type="checkbox"/> This is a SECOND or SUBSEQUENT submission of items concerning a filing under 35 U.S.C. 371. <input checked="" type="checkbox"/> This is an express request to begin national examination procedures (35 U.S.C. 371(f)) at any time rather than delay examination until the expiration of the applicable time limit set in 35 U.S.C. 371(b) and PCT Articles 22 and 39(1). <input checked="" type="checkbox"/> A proper Demand for International Preliminary Examination was made by the 19th month from the earliest claimed priority date. <input checked="" type="checkbox"/> A copy of the International Application as filed (35 U.S.C. 371 (c) (2)) <ol style="list-style-type: none"> <input type="checkbox"/> is transmitted herewith (required only if not transmitted by the International Bureau). <input checked="" type="checkbox"/> has been transmitted by the International Bureau. <input type="checkbox"/> is not required, as the application was filed in the United States Receiving Office (RO/US). <input type="checkbox"/> A translation of the International Application into English (35 U.S.C. 371(c)(2)). <input type="checkbox"/> A copy of the International Search Report (PCT/ISA/210). <input checked="" type="checkbox"/> Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371 (c)(3)) <ol style="list-style-type: none"> <input type="checkbox"/> are transmitted herewith (required only if not transmitted by the International Bureau). <input type="checkbox"/> have been transmitted by the International Bureau. <input type="checkbox"/> have not been made; however, the time limit for making such amendments has NOT expired. <input checked="" type="checkbox"/> have not been made and will not be made. <input type="checkbox"/> A translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371(c)(3)). <input checked="" type="checkbox"/> An oath or declaration of the inventor(s) (35 U.S.C. 371 (c)(4)). <input type="checkbox"/> A copy of the International Preliminary Examination Report (PCT/IPEA/409). <input type="checkbox"/> A translation of the annexes to the International Preliminary Examination Report under PCT Article 36 (35 U.S.C. 371 (c)(5)). 				
Items 13 to 20 below concern document(s) or information included:				
<ol style="list-style-type: none"> <input type="checkbox"/> An Information Disclosure Statement under 37 CFR 1.97 and 1.98. <input type="checkbox"/> An assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included. <input type="checkbox"/> A FIRST preliminary amendment. <input type="checkbox"/> A SECOND or SUBSEQUENT preliminary amendment. <input type="checkbox"/> A substitute specification. <input type="checkbox"/> A change of power of attorney and/or address letter. <input type="checkbox"/> Certificate of Mailing by Express Mail <input type="checkbox"/> Other items or information: 				
Express Mail Label No. EL 813775253				

U.S. APPLICATION NO. (IF KNOWN, SEE PCT/PTO
09/831680INTERNATIONAL APPLICATION NO.
PCT/AU99/00993ATTORNEY'S DOCKET NUMBER
37921-151956

24. The following fees are submitted.:

BASIC NATIONAL FEE (37 CFR 1.492 (a) (1) - (5)) :

<input checked="" type="checkbox"/> Neither international preliminary examination fee (37 CFR 1.482) nor international search fee (37 CFR 1.445(a)(2)) paid to USPTO and International Search Report not prepared by the EPO or JPO	\$1000.00
<input type="checkbox"/> International preliminary examination fee (37 CFR 1.482) not paid to USPTO but International Search Report prepared by the EPO or JPO	\$860.00
<input type="checkbox"/> International preliminary examination fee (37 CFR 1.482) not paid to USPTO but international search fee (37 CFR 1.445(a)(2)) paid to USPTO	\$710.00
<input type="checkbox"/> International preliminary examination fee (37 CFR 1.482) paid to USPTO but all claims did not satisfy provisions of PCT Article 33(1)-(4)	\$690.00
<input type="checkbox"/> International preliminary examination fee (37 CFR 1.482) paid to USPTO and all claims satisfied provisions of PCT Article 33(1)-(4)	\$100.00

CALCULATIONS PTO USE ONLY

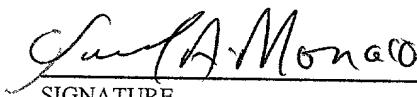
ENTER APPROPRIATE BASIC FEE AMOUNT =			\$1,000.00
Surcharge of \$130.00 for furnishing the oath or declaration later than months from the earliest claimed priority date (37 CFR 1.492 (e)).	<input type="checkbox"/> 20	<input checked="" type="checkbox"/> 30	\$130.00
CLAIMS	NUMBER FILED	NUMBER EXTRA	RATE
Total claims	37 - 20 =	17	x \$18.00 \$306.00
Independent claims	7 - 3 =	4	x \$80.00 \$320.00
Multiple Dependent Claims (check if applicable)		<input checked="" type="checkbox"/>	\$270.00
TOTAL OF ABOVE CALCULATIONS =			\$2,026.00
<input type="checkbox"/> Applicant claims small entity status. (See 37 CFR 1.27). The fees indicated above are reduced by 1/2.			\$0.00
SUBTOTAL =			\$2,026.00
Processing fee of \$130.00 for furnishing the English translation later than months from the earliest claimed priority date (37 CFR 1.492 (f)).	<input type="checkbox"/> 20	<input type="checkbox"/> 30	+ \$0.00
TOTAL NATIONAL FEE =			\$2,026.00
Fee for recording the enclosed assignment (37 CFR 1.21(h)). The assignment must be accompanied by an appropriate cover sheet (37 CFR 3.28, 3.31) (check if applicable).	<input type="checkbox"/>		\$0.00
TOTAL FEES ENCLOSED =			\$2,026.00
		Amount to be: refunded	\$
		charged	\$

- a. A check in the amount of \$2,026.00 to cover the above fees is enclosed.
- b. Please charge my Deposit Account No. _____ in the amount of _____ to cover the above fees. A duplicate copy of this sheet is enclosed.
- c. The Commissioner is hereby authorized to charge any additional fees which may be required, or credit any overpayment to Deposit Account No. 50-0573 A duplicate copy of this sheet is enclosed.
- d. Fees are to be charged to a credit card. **WARNING:** Information on this form may become public. Credit card information should not be included on this form. Provide credit card information and authorization on PTO-2038.

NOTE: Where an appropriate time limit under 37 CFR 1.494 or 1.495 has not been met, a petition to revive (37 CFR 1.137(a) or (b)) must be filed and granted to restore the application to pending status.

SEND ALL CORRESPONDENCE TO:

DANIEL A. MONACO
Drinker Biddle & Reath, LLP
One Logan Square
18th and Cherry Streets
Philadelphia, PA 19103-6996
(215) 988-3312
(215) 988-2757 - Fax



SIGNATURE

DANIEL A. MONACO

NAME

30,480

REGISTRATION NUMBER

May 9, 2001

DATE

09/831686
JCC8 Rec'd PCT/PTO 10 MAY 2001
PATENT

Attorney Docket No.: 37921-151956

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re: Patent application of :
Stephen Locarnini, et al. :
Serial No.: Not Yet Assigned : Group Art Unit:
(International Serial No. PCT/AU99/00993) : Not Yet Assigned
Filed: Concurrently Herewith : Examiner:
(International Filing Date: November 10, 1999) :
For: Biological Compositions, Components : Not Yet Assigned
Thereof and Uses Therefor :
:

PRELIMINARY AMENDMENT

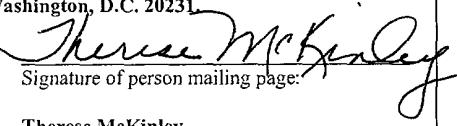
Commissioner for Patents
Box Patent Application
Washington, D.C. 20231

Dear Sir:

Kindly amend the above-identified patent application, without prejudice, as follows.

In the Claims:

Please cancel Claims 14 and 16, without prejudice.

CERTIFICATE OF MAILING UNDER 37 C.F.R. 1.10	
EXPRESS MAIL Mailing Label Number: EL 813775253 US Date of Deposit: May 10, 2001	
I hereby certify that this correspondence, along with any paper referred to as being attached or enclosed, and/or fee, is being deposited with the United States Postal Service, "EXPRESS MAIL-POST OFFICE TO ADDRESSEE" service under 37 CFR 1.10, on the date indicated above, and addressed to: Commissioner for Patents, Washington, D.C. 20231.	
 Signature of person mailing page:	
_____ Therese McKinley Type or print name of person	

Please amend Claims 13 and 15 as follows. A mark-up of the amended claim as required by 37 C.F.R. 1.211(c)(ii) is attached hereto as Appendix A.

13. (Amended) A composition comprising a variant HBV or variant HbsAg according to any one of claims 1 to 12 and one or more pharmaceutically acceptable carriers and/or diluents.

15. (Amended) A method for the treatment or prophylaxis of HBV infection said method comprising administering to a subject an amount of a variant HBV or variant HbsAg according to any one of claims 1 to 12, said amount being effective to induce an immune response to said variant HBV.

REMARKS

Claims 1-13, 15 and 17 are pending in the application. The claims have been amended to reduce claim number.

Respectfully submitted,

STEPHEN LOCARNINI, et al.

BY: 

DANIEL A. MONACO
Drinker, Biddle & Reath, LLP.
One Logan Square
18th and Cherry Streets
Philadelphia, PA 19103-6996
(215) 988-3312
(215) 988-2757 – Fax
Attorney for the Applicants

APPENDIX A - "Marked-up" Versions of Amended Claims as Required Under
37 C.F.R. 1.121(c)(1)(ii)

13. (Amended) A composition comprising a variant HBV or variant HbsAg according to any one of claims 1 to 12 [or a recombinant or derivative form or its chemical equivalent] and one or more pharmaceutically acceptable carriers and/or diluents.

15. (Amended) A method for the treatment or prophylaxis of HBV infection said method comprising administering to a subject an amount of a variant HBV or variant HbsAg according to any one of claims 1 to 12 [or a composition according to claim 13 or 14], said amount being effective to induce an immune response to said variant HBV.

BIOLOGICAL COMPOSITIONS, COMPONENTS THEREOF AND USES THEREFOR

5 FIELD OF THE INVENTION

The present invention relates generally to an isolated Hepatitis B virus (HBV) with a surface component exhibiting an altered immunological profile relative to a reference HBV. A reference HBV is considered herein to comprise a composite or consensus nucleotide or amino acid sequence from HBV genotypes A through F. The isolated HBV of the present invention is considered herein to be a HBV variant relative to the reference HBV. The altered immunological profile renders the HBV variants of the present invention less susceptible to vaccines directed to the surface component. The HBV variants of the present invention generally arise from selective pressure following one or both of anti-HBV chemical therapy and in particular chemical therapy aimed at disrupting HBV polymerase activity or function and/or following immune pressure directed to the surface component. Immune pressure may result from natural exposure to HBV or following vaccination with an avirulent or attenuated HBV or with a component of an HBV. The present invention further provides a recombinant polypeptide and derivatives and chemical equivalents thereof corresponding to the surface component of the HBV variants. The HBV variants and recombinant polypeptides and their derivatives and chemical equivalents of the present invention are useful in biological compositions capable of inducing a neutralizing immune response to the HBV variant.

25 BACKGROUND OF THE INVENTION

Bibliographic details of the publications numerically referred to in this specification are collected at the end of the description.

30 The rapidly increasing sophistication of recombinant DNA technology is greatly

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facilitating advances in the medical and allied health fields. This is particularly the case with the generation of recombinant vaccines and therapeutic compositions. Recombinant technology is providing the means to generate recombinant components of vaccines as well as providing genetic bases for screening or identifying useful components for therapeutic compositions.

Hepatitis B virus (HBV) can cause debilitating disease conditions ranging from subclinical infection to chronic active and fulminant hepatitis and can lead to acute liver failure.

10 The HBV genome comprises a series of overlapping genes in a circular, partially double-stranded DNA molecule (1) [see also Figure 1]. For example, the gene encoding DNA polymerase overlaps the viral envelope genes, Pre-S1, Pre-S2 and S and partially overlaps the X and core genes. The HBV envelope comprises small, middle and large HBV surface proteins. The large protein component is generally referred to as the HBV surface antigen 15 (HBsAg) and is encoded by the S gene sequence. The Pre-S1 and Pre-S2 gene sequences encode the other envelope components (2).

The HBsAg comprises an antigenic region referred to as the "a" determinant (3). The "a" determinant is complex, conformational and dependent upon disulphide bonding among 20 highly conserved cysteine residues. Genetic variation leading to changes in the "a" determinant has been implicated in mutants of HBV which "escape" the immunological response generated to conventional vaccines (4-8). One particularly common mutation is a glycine (G) to arginine (R) substitution at amino acid position 145 (G145R) of HBsAg. This mutation affects the "a" epitope region.

25

The increasing reliance on chemical and immunological intervention in treating or preventing HBV infection is resulting in greater selective pressure for the emergence of variants of HBV which are resistant to the interventionist therapy. Such variants are referred to as "escape" mutants.

30

There is a need to foreshadow potential vaccine escape variants of HBV such that biological compositions can be quickly prepared for use as vaccines directed against the modified virus or its altered antigenic components.

5 SUMMARY OF THE INVENTION

Specific mutations in amino acid sequence are represented herein as "Xaa₁nXaa₂" where Xaa₁ is the original amino acid residue before mutation, n is the residue number and Xaa₂ is the mutant amino acid. The abbreviation "Xaa" may be the three letter or single letter 10 amino acid code. A mutation in single letter code is represented, for example, by X₁nX₂ where X₁ and X₂ are the same as Xaa₁ and Xaa₂, respectively. The amino acid residues for Hepatitis B virus DNA polymerase are numbered with the residue methionine in the motif Tyr Met Asp Asp (YMDD) being residue number 550.

15 The reference HBV is considered herein to comprise a composite or consensus nucleotide or amino acid sequence from HBV genotypes A through F.

One aspect of the present invention provides a variant HBV comprising a surface component exhibiting an altered immunological profile compared to a reference HBV.

20 Another aspect of the present invention is directed to a variant HBV comprising a surface antigen having an amino acid sequence with a single or multiple amino acid substitution, addition and/or deletion or a truncation compared to a surface antigen from a reference HBV and wherein an antibody generated to the reference surface antigen exhibits reduced 25 capacity for neutralizing said HBV variant.

Yet another aspect of the present invention provides an HBV variant comprising a surface antigen having an amino acid sequence with a single or multiple amino acid substitution, addition and/or deletion or truncation compared to the amino acid sequence set forth in 30 Formula I and wherein the surface antigen of the variant HBV exhibits an altered

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immunological profile compared to the surface antigen defined by Formula I and wherein the variant HBV is selected for by a nucleoside analogue of the HBV DNA polymerase.

Still another aspect of the present invention is directed to an HBV variant comprising a
5 surface antigen having an amino acid sequence with a single or multiple amino acid substitution, addition and/or deletion or truncation compared to the amino acid sequence set forth in Formula I and wherein the surface antigen of the variant HBV exhibits an altered immunological profile compared to the surface antigen defined by Formula I and wherein the variant HBV is selected for following immunological therapy directed against
10 the surface antigen as defined in Formula I.

Even still another aspect of the present invention provides an HBV variant comprising a nucleotide sequence comprising a single or multiple nucleotide substitution, addition and/or deletion to the nucleotide sequence set forth in Formula III and which HBV variant
15 has a surface antigen exhibiting an altered immunological profile relative to a surface antigen defined by Formula I.

Another aspect of the present invention provides an isolated HBsAg or a recombinant form thereof or derivative or chemical equivalent thereof.
20

Yet another aspect of the present invention is directed to an isolated variant HBsAg or a recombinant or derivative form thereof or a chemical equivalent thereof wherein said HBsAg or its recombinant or derivative form or its chemical equivalent exhibits an altered immunological profile compared to an HBsAg from a reference HBV.
25

Still yet another aspect of the present invention provides an isolated variant HBsAg or a recombinant or derivative form thereof or a chemical equivalent thereof wherein said HBsAg or its recombinant or derivative form or its chemical equivalent comprises an amino acid sequence with a single or multiple amino acid substitution, addition and/or
30 deletion or a truncation compared to an HBsAg from a reference HBV and wherein a

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neutralising antibody directed to a reference HBV exhibits no or reduced neutralising activity to an HBV carrying said variant HBsAg.

Another aspect of the present invention contemplates a biological composition comprising a
5 variant HBV or an HBsAg from said variant HBV or a recombinant or derivative form thereof or its chemical equivalent.

BRIEF DESCRIPTION OF THE FIGURES

Figure 1 is a diagrammatic representation showing overlapping genome of HBV.

5

Figure 2 is a representation of the amino acid consensus sequence from HBV DNA polymerase proteins encompassing regions which are conserved in the RNA polymerase protein. These regions are shown as domains A-E and are underlined. In the consensus sequence the M in the YMDD motif is designated as amino acid number 550. The amino acids which are subject to mutation during 3TC and/or FCV treatment are shown in bold. An asterisk (*) indicates greater than three amino acid possibilities at this position of the consensus sequence. The HBsAg major hydrophilic region containing the neutralisation domain is indicated by a double line and the polymerase mutations which alter the HBsAg are indicated in italics.

15

Figure 3 is a representation of the nucleotide sequence from various strains of HBV encoding the surface antigen. The amino acid sequence of the surface antigen beginning at amino acid 108 is shown above the nucleotide sequence.

20 **Figure 4** is a graphical representation showing HBsAg binding assay with wild-type (i.e. reference HBV) and various mutants (1, mock; 2, wild-type; 3, F512L; 4, V519L; 5, M550I; 6, S565P; 7, double mutant L256M + M550V; 8, triple mutant V519L + L526M + M550V; 9, W499Q).

25

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

The present invention is predicated in part on the identification of HBV variants exhibiting an altered immunological profile with respect to a surface component relative to a reference standard. The term "variant" is used in its broadest context and includes mutants such as vaccine escape mutants, derivatives, modified forms of and altered forms of an HBV relative to a reference HBV. A variant generally contains a single or multiple nucleotide substitution, addition and/or deletion or a truncation mutation in the viral genome and a corresponding single or multiple amino acid substitution, addition and/or 10 deletion or truncation in a viral peptide, polypeptide or protein.

A preferred variant in accordance with the present invention with an altered immunological profile is one which would substantially not be affected by a neutralizing immune response directed to a conventional HBV vaccine such as a vaccine comprising a reference HBV or a surface component thereof. The expression "substantially not affected" includes reduced susceptibility to the immune response generated by a vaccine. Reduced susceptibility may also be conveniently determined by reduced susceptibility to chemical agents such as nucleoside analogues which target HBV DNA polymerase. Due to the overlapping nature of reading frames for DNA polymerase and certain viral surface 20 components, an altered surface component may have a corresponding alteration in the DNA polymerase.

The preferred surface component of the HBV of the present invention is the HBV surface antigen (HBsAg). It is proposed in accordance with the present invention that the HBsAg 25 of the HBV variants exhibit an altered immune profile relative to an HBsAg from a reference HBV. For the purposes of the present invention, a reference HBV conveniently comprises an HBsAg with an amino acid sequence substantially as set forth by Norder *et al.* (9) which encompasses all known genotypes of HBV (currently A through F). The amino acid sequence of an HBsAg and which is considered to define a reference HBV is 30 set forth below in Formula I:

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FORMULA I

M X₁ X₂ X₃ X₄ S G X₅ L X₆ P L X₇ V L Q A X₈ X₉ F X₁₀ L T X₁₁ I X₁₂ X₁₃ I P
X₁₄ S L X₁₅ S W W T S L N F L G X₁₆ X₁₇ X₁₈ X₁₉ C X₂₀ G X₂₁ N X₂₂ Q S
5 X₂₃ X₂₄ S X₂₅ H X₂₆ P X₂₇ X₂₈ C P P X₂₉ C X₃₀ G Y R W M C L X₃₁ R F I I F
L X₃₂ I L L L C L I F L L V L L D X₃₃ Q G M L X₃₄ V C P L X₃₅ P X₃₆ X₃₇ X₃₈
T T S X₃₉ X₄₀ X₄₁ C X₄₂ T C X₄₃ X₄₄ X₄₅ X₄₆ Q G X₄₇ S X₄₈ X₄₉ P X₅₀ X₅₁ C
C X₅₂ K P X₅₃ X₅₄ G N C T C I P I P S X₅₅ W A X₅₆ X₅₇ X₅₈ X₅₉ L W E X₆₀
X₆₁ S X₆₂ R X₆₃ S W L X₆₄ L L X₆₅ X₆₆ F V Q X₆₇ X₆₈ X₆₉ X₇₀ L X₇₁ P X₇₂ V W
10 X₇₃ X₇₄ X₇₅ I W X₇₆ X₇₇ W X₇₈ W X₇₉ P X₈₀ X₈₁ X₈₂ X₈₃ I X₈₄ X₈₅ P F X₈₆ P L
L P I F X₈₇ X₈₈ L X₈₉ X₉₀ X₉₁ I

wherein:

X₁ is E or G or D;
15 X₂ is N or S or K;
X₃ is I or T;
X₄ is T or A;
X₅ is F or L;
X₆ is G or R;
20 X₇ is L or R;
X₈ is G or V;
X₉ is F or C;
X₁₀ is L or S or W;
X₁₁ is R or K;
25 X₁₂ is L or R;
X₁₃ is T or K;
X₁₄ is Q or K;

X₁₅ is D or H;
X₁₆ is G or E or A;
X₁₇ is S or A or V or T or L;
X₁₈ is P or T;
5 X₁₉ is V or R or T or K or G;
X₂₀ is L or P;
X₂₁ is Q or L or K;
X₂₂ is S or L;
X₂₃ is P or Q;
10 X₂₄ is T or I;
X₂₅ is N or S;
X₂₆ is S or L;
X₂₇ is T or I;
X₂₈ is S or C;
15 X₂₉ is I or T;
X₃₀ is P or A;
X₃₁ is R or Q;
X₃₂ is F or C;
X₃₃ is Y or C;
20 X₃₄ is P or H or S;
X₃₅ is I or L;
X₃₆ is G or R;
X₃₇ is S or T;
X₃₈ is T or S;
25 X₃₉ is T or V or A;
X₄₀ is G or E or Q;
X₄₁ is P or A or S;

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X_{42} is K or R;
 X_{43} is T or M;
 X_{44} is T or I or S or A;
 X_{45} is P or T or A or I or L;
5 X_{46} is A or V;
 X_{47} is N or T;
 X_{48} is M or K or L;
 X_{49} is F or Y or I;
 X_{50} is S or Y;
10 X_{51} is C or S;
 X_{52} is T or I or S;
 X_{53} is T or S;
 X_{54} is D or A;
 X_{55} is S or T;
15 X_{56} is F or L;
 X_{57} is A or G or V;
 X_{58} is K or R or T;
 X_{59} is Y or F;
 X_{60} is W or G;
20 X_{61} is A or G;
 X_{62} is V or A;
 X_{63} is F or L;
 X_{64} is S or N;
 X_{65} is V or A;
25 X_{66} is P or Q;
 X_{67} is W or C or S;
 X_{68} is F or C;

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X₆₉ is V or D or A;
X₇₀ is G or E;
X₇₁ is S or F;
X₇₂ is T or I;
5 X₇₃ is L or P;
X₇₄ is S or L;
X₇₅ is A or V;
X₇₆ is M or I;
X₇₇ is M or I;
10 X₇₈ is Y or F;
X₇₉ is G or E;
X₈₀ is S or N or K;
X₈₁ is L or Q;
X₈₂ is Y or F or H or C;
15 X₈₃ is S or G or N or D or T;
X₈₄ is V or L;
X₈₅ is S or N;
X₈₆ is I or M or L;
X₈₇ is F or C;
20 X₈₈ is C or Y;
X₈₉ is W or R;
X₉₀ is V or A; and
X₉₁ is Y or I or S.

25 Accordingly, one aspect of the present invention provides a variant HBV comprising a surface component exhibiting an altered immunological profile compared to a reference HBV.

More particularly, the present invention is directed to a variant HBV comprising a surface antigen having an amino acid sequence with a single or multiple amino acid substitution, addition and/or deletion or a truncation compared to a surface antigen from a reference HBV and wherein an antibody generated to the reference surface antigen exhibits reduced capacity for neutralizing said HBV variant.

The amino acid sequence of the HBsAg of the reference HBV is as set forth in Formula I above.

10 The HBV variant of the present invention is also referred to herein as an "escape" mutant since it is substantially incapable of being adversely effected by chemical therapy directed against the HBV polymerase or vaccine therapy directed against the surface antigen. The term "escape" mutant also encompasses reduced susceptibility to chemical or vaccine therapy directed to the reference HBV.

15

The HBV variant of the present invention is also preferably in isolated form. An isolated HBV includes reference to a biologically pure form of the virus. The term "isolated" means the virus has undergone at least one purification or isolation step away from non-viral components. Preferably, the viral preparation comprises at least about 10%, more 20 preferably at least about 20%, still more preferably at least about 30%, even more preferably at least about 40%, yet more preferably at least about 50% or greater of HBV variant relative to the non-viral components as measured by viral infectivity, immunological interactivity, DNA polymerase activity, molecular weight, carbohydrate content or other suitable means.

25

The preferred variants of the present invention are obtained following selective pressure. The preferred selective pressure is chemical pressure (e.g. *via* nucleoside analogues) directed to the HBV DNA polymerase which selects for a mutation in the gene encoding HBV DNA polymerase and a corresponding mutation in the gene encoding HBsAg. This 30 is due to the overlapping open reading frames for HBV DNA polymerase and HBsAg. A

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mutation in one or more nucleotides encoding HBV DNA polymerase may have an effect on the nucleotide sequence encoding HBsAg. The present invention also extends to changes in the HBsAg following immunological selection based on vaccines comprising HBsAg or a derivative thereof or an HBV comprising same and wherein the HBsAg 5 comprises an amino acid sequence substantially as set forth in Formula I.

Accordingly, another aspect of the present invention provides an HBV variant comprising a surface antigen having an amino acid sequence with a single or multiple amino acid substitution, addition and/or deletion or truncation compared to the amino acid sequence 10 set forth in Formula I and wherein the surface antigen of the variant HBV exhibits an altered immunological profile compared to the surface antigen defined by Formula I and wherein the variant HBV is selected for by a nucleoside analogue of the HBV DNA polymerase.

15 In a related embodiment the present invention is directed to an HBV variant comprising a surface antigen having an amino acid sequence with a single or multiple amino acid substitution, addition and/or deletion or truncation compared to the amino acid sequence set forth in Formula I and wherein the surface antigen of the variant HBV exhibits an altered immunological profile compared to the surface antigen defined by Formula I and 20 wherein the variant HBV is selected for following immunological therapy directed against the surface antigen as defined in Formula I.

Reference to an altered immunological profile in accordance with the present invention in relation to the surface antigen includes reference to an altered humoral or T cell response.

25 Examples of an altered immunological profile include altered specificity to antibodies, altered amino acid sequences of an epitope or within the "a" determinant, an altered capacity to induce proliferation of T cells primed to an HBsAg from a reference HBV. Preferably, the altered immunological profile means that neutralising antibodies which are capable of substantially neutralising or otherwise reducing serum or blood levels of the 30 reference HBV are substantially incapable of or exhibit reduced capacity to neutralise

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and/or clear the variant HBV.

A viral variant may, in accordance with a preferred aspect of the present invention, carry a mutation only in the DNA polymerase or the surface antigen or may carry a mutation in both molecules. The term "mutation" is to be read in its broadest context and includes a silent mutation not substantially affecting the normal function of the DNA polymerase or surface antigen or may be an active mutation having the effect of selection of nucleoside analogue resistance or a vaccine escape mutant phenotype. Where multiple mutations occur in accordance with the present invention or where multiple phenotypes result from a single mutation, at least one mutation must be active or the virus must exhibit at least one altered phenotype such as nucleoside analogue resistance or reduced immunological interactivity to the surface antigen of a reference HBV.

The present invention extends to any novel mutant or novel use of a mutant of the HBsAg carrying a single or multiple substitution, addition and/or deletion or truncation in the amino acid sequence of HBsAg as compared to the amino acid sequence set forth in Formula I. In an alternative yet related embodiment, the present invention extends to any single or multiple amino acid substitution, addition and/or deletion or truncation in the amino acid sequence of HBsAg relative to the amino acid sequence set forth in Formula I as defined by a single or multiple amino acid substitution, addition and/or deletion to the catalytic region of the HBV DNA polymerase set forth below in Formula II:

FORMULA II

S Z₁ L S W L S L D V S A A F Y H Z₂ P L H P A A M P H L L Z₃ G S S G
L Z₄ R Y V A R L S S Z₅ S Z₆ Z₇ X N Z₈ Q Z₉ Z₁₀ X X X Z₁₁ L H Z₁₂ Z₁₃ C S
R Z₁₄ L Y V S L Z₁₅ L L Y Z₁₆ T Z₁₇ G Z₁₈ K L H L Z₁₉ Z₂₀ H P I Z₂₁ L G F R
K Z₂₂ P M G Z₂₃ G L S P F L L A Q F T S A I Z₂₄ Z₂₅ Z₂₆ Z₂₇ Z₂₈ R A F Z₂₉
H C Z₃₀ Z₃₁ F Z₃₂ Y M* D D Z₃₃ V L G A Z₃₄ Z₃₅ Z₃₆ Z₃₇ H Z₃₈ E Z₃₉ L Z₄₀ Z₄₁

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$Z_{42} Z_{43} Z_{44} Z_{45} Z_{46} L L Z_{47} Z_{48} G I H L N P Z_{49} K T K R W G Y S L N F M G$
 $Y Z_{50} I G$

wherein:

- 5 X is any amino acid;
- Z_1 is N or D;
- Z_2 is I or P;
- Z_3 is I or V;
- Z_4 is S or D;
- 10 Z_5 is T or N;
- Z_6 is R or N;
- Z_7 is N or I;
- Z_8 is N or Y or H;
- Z_9 is H or Y;
- 15 Z_{10} is G or R;
- Z_{11} is D or N;
- Z_{12} is D or N;
- Z_{13} is S or Y;
- Z_{14} is N or Q;
- 20 Z_{15} is L or M;
- Z_{16} is K or Q;
- Z_{17} is Y or F;
- Z_{18} is R or W;
- Z_{19} is Y or L;
- 25 Z_{20} is S or A;
- Z_{21} is I or V;
- Z_{22} is I or L;

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Z_{23} is V or G;
 Z_{24} is C or L;
 Z_{25} is A or S;
 Z_{26} is V or M;
5 Z_{27} is V or T;
 Z_{28} is R or C;
 Z_{29} is F or P;
 Z_{30} is L or V;
 Z_{31} is A or V;
10 Z_{32} is S or A;
 Z_{33} is V or L or M;
 Z_{34} is K or R;
 Z_{35} is S or T;
 Z_{36} is V or G;
15 Z_{37} is Q or E;
 Z_{38} is L or S or R;
 Z_{39} is S or F;
 Z_{40} is F or Y;
 Z_{41} is T or A;
20 Z_{42} is A or S;
 Z_{43} is V or I;
 Z_{44} is T or C;
 Z_{45} is N or S;
 Z_{46} is F or V;
25 Z_{47} is S or D;
 Z_{48} is L or V;
 Z_{49} is N or Q;

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Z_{50} is V or I; and

M* is amino acid 550.

Preferred mutations in the amino acid sequence of HBsAg are amino acid substitutions,

5 deletions and/or additions or truncations in amino acids 1-10, 5-15, 10-20, 15-25, 20-30, 25-
35, 30-40, 35-45, 40-50, 45-55, 50-60, 55-65, 60-70, 65-75, 70-80, 75-85, 80-90, 85-95, 90-
100, 95-105, 100-110, 105-115, 110-120, 115-125, 120-130, 125-135, 130-140, 135-145,
140-150, 145-155, 150-160, 155-165, 160-170, 165-175, 170-180, 175-185, 180-190, 185-
195, 190-200, 195-205, 200-210, 205-215, 210-220, 215-225, 220-226 (referring to the
10 numbering of Formula I) of HBsAg. Particularly useful mutations are G112R, T123P
Y/F134S, D144E, G145R, A157D, E164D, F170L, M195I, W196L, S196W, W196 STOP,
M198I, W199S, S204T and S210R. The term "stop" means a stop codon.

Even more preferred mutations are D144E, G145R, A157D, E164D, M195I, W196L,

15 S196W, W196 STOP, M198I, W199S and S210R.

The HBsAg mutations of the present invention may also be defined in terms of a corresponding mutation in the HBV DNA polymerase. A mutation in the HBV DNA polymerase may be in amino acids 421-431, 426-436, 431-441, 436-446, 441-451, 446-456,
20 451-461, 456-466, 461-471, 466-476, 471-481, 476-486, 481-491, 486-496, 491-501, 496-
506, 501-511, 506-516, 511-521, 516-526, 521-531, 526-536, 531-541, 536-546, 541-551,
546-556, 551-561, 556-566, 561-571, 566-576, 571-581, 576-586, 581-591, 586-596, 591-
601, 596-601 (referring to number of Formula II).

25 Preferred HBV DNA polymerase mutations include Q476, N480G, N485K, K495R, R499O,
G499E, W499Q, F112L, I151L, V519L, L526M, M550V, M550I, V553I, S565P. Useful
multiple mutants include L526M/M550I, L526M/M550V, V519L/L526M/M550V and
V519L/L526M/M550I.

30 The altered HBsAg molecules of the HBV variants of the present invention may also be

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defined at the nucleotide level. The nucleotide sequence encoding the HBsAg from a reference HBV is set forth below in Formula III:

FORMULA III

5

A C N₁ A A A C C T N₂ N₃ G G A N₄ G G A A A N₅ T G C A C N₆ T G T A
T T C C C A T C C C A T C N₇ T C N₈ T G G G C T T C G N₉ A A N₁₀
A T N₁₁ C C T A T G G G A G N₁₂ G G G C C T C A G N₁₃ C C G T T T
C T C N₁₄ T G G C T C A G T T A C T A G T G C C A T T G T T C A
10 G T G G T T C G N₁₅ A G G G C T T C C C C C A C T G T N₁₆ T G G
C T T C A G N₁₇ T A T A T G G A T G A T G T G G T N₁₈ T T G G G G
G C C A A G T C T G T A C A N₁₉ C A T C N₂₀ T G A G T C C C T T T
N₂₁ T N₂₂ C C N₂₃ C T N₂₄ T T A C C A A T T T C T T N₂₅ T G T C T N₂₆
T G G G N₂₇ A T A C A T T

15

wherein:

N₁ is A or C;

N₂ is T or A;

N₃ is C or T;

20 N₄ is C or T;

N₅ is C or T;

N₆ is C or T;

N₇ is A or G;

N₈ is T or C;

25 N₉ is C or G;

N₁₀ is G or A;

N₁₁ is T or A;

N₁₂ is T or G;

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N₁₃ is T or C;
N₁₄ is C or T;
N₁₅ is T or C;
N₁₆ is T or C;
5 N₁₇ is T or C;
N₁₈ is A or T;
N₁₉ is A or G;
N₂₀ is T or G;
N₂₁ is A or T;
10 N₂₂ is A or G;
N₂₃ is T or G;
N₂₄ is A or G;
N₂₅ is T or C;
N₂₆ is T or C; and
15 N₂₇ is T or C.

The present invention extends to nucleotide sequences which exhibit at least about 60% nucleotide sequence identity to Formula III or is a sequence capable of hybridising thereto under low stringency conditions at 42 °C and which encode an HBsAg with an altered 20 immunological profile relative to an HBsAg from a reference HBV.

Accordingly, another aspect of the present invention provides an HBV variant comprising a nucleotide sequence comprising a single or multiple nucleotide substitution, addition and/or deletion to the nucleotide sequence set forth in Formula III and which HBV variant has a 25 surface antigen exhibiting an altered immunological profile relative to a surface antigen defined by Formula I.

Preferably, the HBV variant comprises a nucleotide sequence having at least about 80% identity to the nucleotide sequence set forth in Formula III or is capable of hybridising

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thereto under medium stringency conditions at 42 °C. Preferably, the percentage identity is at least about 85%, at least about 90%, at least about 95%, but less than 100% relative to the nucleotide sequence set forth in Formula III.

5 The term "similarity" as used herein includes exact identity between compared sequences at the nucleotide or amino acid level. Where there is non-identity at the nucleotide level, "similarity" includes differences between sequences which result in different amino acids that are nevertheless related to each other at the structural, functional, biochemical and/or conformational levels. Where there is non-identity at the amino acid level, "similarity" 10 includes amino acids that are nevertheless related to each other at the structural, functional, biochemical and/or conformational levels. In a particularly preferred embodiment, nucleotide and sequence comparisons are made at the level of identity rather than similarity. Any number of programs are available to compare nucleotide and amino acid sequences. Preferred programs have regard to an appropriate alignment. One such program is Gap 15 which considers all possible alignment and gap positions and creates an alignment with the largest number of matched bases and the fewest gaps. Gap uses the alignment method of Needleman and Wunsch (10). Gap reads a scoring matrix that contains values for every possible GCG symbol match. GAP is available on ANGIS (Australian National Genomic Information Service) at website <http://mell.angis.org.au..>

20

Reference herein to a low stringency at 42 °C includes and encompasses from at least about 0% v/v to at least about 15% v/v formamide and from at least about 1M to at least about 2M salt for hybridisation, and at least about 1M to at least about 2M salt for washing conditions. Alternative stringency conditions may be applied where necessary, such as medium

25 stringency, which includes and encompasses from at least about 16% v/v to at least about 30% v/v formamide and from at least about 0.5M to at least about 0.9M salt for hybridisation, and at least about 0.5M to at least about 0.9M salt for washing conditions, or high stringency, which includes and encompasses from at least about 31% v/v to at least about 50% v/v formamide and from at least about 0.01M to at least about 0.15M salt for 30 hybridisation, and at least about 0.01M to at least about 0.15M salt for washing conditions. In general, washing is carried out $T_m = 69.3 + 0.41 (G+C)\%$ [11]. However, the T_m of a

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duplex DNA decreases by 1°C with every increase of 1% in the number of mismatch base pairs (12).

The present invention further extends to an isolated surface component from the HBV variants herein described. More particularly, the present invention provides an isolated HBsAg or a recombinant form thereof or derivative or chemical equivalent thereof. The isolated surface component and, more particularly, isolated HBsAg or its recombinant, derivative or chemical equivalents are useful in the development of biological compositions such as vaccine formulations.

10

Accordingly, another aspect of the present invention is directed to an isolated variant HBsAg or a recombinant or derivative form thereof or a chemical equivalent thereof wherein said HBsAg or its recombinant or derivative form or its chemical equivalent exhibits an altered immunological profile compared to an HBsAg from a reference HBV.

15

More particularly, the present invention provides an isolated variant HBsAg or a recombinant or derivative form thereof or a chemical equivalent thereof wherein said HBsAg or its recombinant or derivative form or its chemical equivalent comprises an amino acid sequence with a single or multiple amino acid substitution, addition and/or deletion or a 20 truncation compared to an HBsAg from a reference HBV and wherein a neutralising antibody directed to a reference HBV exhibits no or reduced neutralising activity to an HBV carrying said variant HBsAg.

The term "isolated" means the same as it does in relation to an isolated HBV variant.

25

The reference HBV is conveniently defined herein as comprising an HBsAg with an amino acid sequence as set forth in Formula I or as indirectly defined by the amino acid sequence for HBV DNA polymerase set forth in Formula II or by the nucleotide sequence set forth in Formula III encoding an HBsAg.

30

As stated above, the present invention extends to derivatives and chemical equivalents (i.e.

analogues) of the HBV surface component and in particular HBsAg. Derivatives include single or multiple amino acid substitutions, additions and/or deletions to the HBsAg molecule. "Additions" to amino acid sequences include fusions with other peptides, polypeptides or proteins or fusions to nucleotide sequences including fusions to other viral components.

Analogues of the variant HBsAg contemplated herein include, but are not limited to, modification to side chains, incorporating of unnatural amino acids and/or their derivatives during peptide, polypeptide or protein synthesis and the use of crosslinkers and other methods which impose conformational constraints on the proteinaceous molecule or their analogues. These types of modifications are useful in stabilizing the immunointeractive molecules for use in diagnostic assays or in therapeutic protocols.

Examples of side chain modifications contemplated by the present invention include modifications of amino groups such as by reductive alkylation by reaction with an aldehyde followed by reduction with NaBH₄; amidination with methylacetimidate; acylation with acetic anhydride; carbamoylation of amino groups with cyanate; trinitrobenzylolation of amino groups with 2, 4, 6-trinitrobenzene sulphonic acid (TNBS); acylation of amino groups with succinic anhydride and tetrahydrophthalic anhydride; and pyridoxylation of lysine with pyridoxal-5-phosphate followed by reduction with NaBH₄.

The guanidine group of arginine residues may be modified by the formation of heterocyclic condensation products with reagents such as 2,3-butanedione, phenylglyoxal and glyoxal.

The carboxyl group may be modified by carbodiimide activation *via* O-acylisourea formation followed by subsequent derivitisation, for example, to a corresponding amide.

Sulphydryl groups may be modified by methods such as carboxymethylation with iodoacetic acid or iodoacetamide; performic acid oxidation to cysteic acid; formation of a mixed disulphides with other thiol compounds; reaction with maleimide, maleic anhydride or other substituted maleimide; formation of mercurial derivatives using 4-chloromercuribenzoate,

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4-chloromercuriphenylsulphonic acid, phenylmercury chloride, 2-chloromercuri-4-nitrophenol and other mercurials; carbamoylation with cyanate at alkaline pH.

Tryptophan residues may be modified by, for example, oxidation with N-bromosuccinimide or alkylation of the indole ring with 2-hydroxy-5-nitrobenzyl bromide or sulphenyl halides.

Tyrosine residues on the other hand, may be altered by nitration with tetranitromethane to form a 3-nitrotyrosine derivative.

Modification of the imidazole ring of a histidine residue may be accomplished by alkylation with iodoacetic acid derivatives or N-carbethoxylation with diethylpyrocarbonate.

Examples of incorporating unnatural amino acids and derivatives during peptide synthesis include, but are not limited to, use of norleucine, 4-amino butyric acid, 4-amino-3-hydroxy-5-phenylpentanoic acid, 6-aminohexanoic acid, t-butylglycine, norvaline, phenylglycine, ornithine, sarcosine, 4-amino-3-hydroxy-6-methylheptanoic acid, 2-thienyl alanine and/or D-isomers of amino acids. A list of unnatural amino acid, contemplated herein is shown below in Table 1. The inclusion of such unnatural amino acids or other derivations described herein may assist in stabilising the molecule in a vaccine composition.

TABLE 1

Non-conventional amino acid	Code	Non-conventional amino acid	Code
5 α -aminobutyric acid	Abu	L-N-methylalanine	Nmala
α -amino- α -methylbutyrate	Mgabu	L-N-methylarginine	Nmarg
aminocyclopropane- carboxylate	Cpro	L-N-methyleasparagine	Nmasn
		L-N-methyleaspartic acid	Nmasp
10 aminoisobutyric acid	Aib	L-N-methylcysteine	Nmcys
aminonorbornyl- carboxylate	Norb	L-N-methylglutamine	Nmgln
		L-N-methylglutamic acid	Nmglu
cyclohexylalanine		Chexa L-N-methylhistidine	Nmhis
cyclopentylalanine	Cpen	L-N-methylisoleucine	Nmile
15 D-alanine	Dal	L-N-methylleucine	Nmleu
D-arginine	Darg	L-N-methyllysine	Nmlys
D-aspartic acid	Dasp	L-N-methylmethionine	Nmmet
D-cysteine	Dcys	L-N-methylnorleucine	Nmnle
D-glutamine	Dgln	L-N-methylnorvaline	Nmnva
20 D-glutamic acid	Dglu	L-N-methylornithine	Nmom
D-histidine	Dhis	L-N-methylphenylalanine	Nmphe
D-isoleucine	Dile	L-N-methylproline	Nmpro
D-leucine	Dleu	L-N-methylserine	Nmser
D-lysine	Dlys	L-N-methylthreonine	Nmthr
25 D-methionine	Dmet	L-N-methyltryptophan	Nmtrp
D-ornithine	Dorn	L-N-methyltyrosine	Nmtyr
D-phenylalanine	Dphe	L-N-methylvaline	Nmval
D-proline	Dpro	L-N-methylethylglycine	Nmetg
D-serine	Dser	L-N-methyl-t-butylglycine	Nmtbug
30 D-threonine	Dthr	L-norleucine	Nle
D-tryptophan	Dtrp	L-norvaline	Nva

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D-tyrosine	Dtyr	α -methyl-aminoisobutyrate	Maib
D-valine	Dval	α -methyl- γ -aminobutyrate	Mgabu
D- α -methylalanine	Dmala	α -methylcyclohexylalanine	Mchexa
D- α -methylarginine	Dmarg	α -methylcyclopentylalanine	Mcpen
5 D- α -methylasparagine	Dmasn	α -methyl- α -naphthylalanine	Manap
D- α -methylaspartate	Dmasp	α -methylpenicillamine	Mpen
D- α -methylcysteine	Dmcys	N-(4-aminobutyl)glycine	Nglu
D- α -methylglutamine	Dmgln	N-(2-aminoethyl)glycine	Naeg
D- α -methylhistidine	Dmhis	N-(3-aminopropyl)glycine	Norn
10 D- α -methylisoleucine	Dmile	N-amino- α -methylbutyrate	Nmaabu
D- α -methylleucine	Dmleu	α -naphthylalanine	Anap
D- α -methyllysine	Dmlys	N-benzylglycine	Nphe
D- α -methylmethionine	Dmmet	N-(2-carbamylethyl)glycine	Ngln
D- α -methylornithine	Dmorn	N-(carbamylmethyl)glycine	Nasn
15 D- α -methylphenylalanine	Dmphe	N-(2-carboxyethyl)glycine	Nglu
D- α -methylproline	Dmpro	N-(carboxymethyl)glycine	Nasp
D- α -methylserine	Dmser	N-cyclobutylglycine	Ncbut
D- α -methylthreonine	Dmthr	N-cycloheptylglycine	Nchep
D- α -methyltryptophan	Dmtrp	N-cyclohexylglycine	Nchex
20 D- α -methyltyrosine	Dmty	N-cyclodecylglycine	Ncdec
D- α -methylvaline	Dmval	N-cyclododecylglycine	Ncdod
D-N-methylalanine	Dnmala	N-cyclooctylglycine	Neoct
D-N-methylarginine	Dnmarg	N-cyclopropylglycine	Ncpro
D-N-methylasparagine	Dnmasn	N-cycloundecylglycine	Ncund
25 D-N-methylaspartate	Dnmasp	N-(2,2-diphenylethyl)glycine	Nbhm
D-N-methylcysteine	Dnmcys	N-(3,3-diphenylpropyl)glycine	Nbhe
D-N-methylglutamine	Dnmgln	N-(3-guanidinopropyl)glycine	Narg
D-N-methylglutamate	Dnmglu	N-(1-hydroxyethyl)glycine	Nthr
D-N-methylhistidine	Dnmhis	N-(hydroxyethyl)glycine	Nser
30 D-N-methylisoleucine	Dnmile	N-(imidazolylethyl)glycine	Nhis
D-N-methylleucine	Dnmleu	N-(3-indolylethyl)glycine	Nhtrp

D-N-methyllysine	Dnmlys	N-methyl- γ -aminobutyrate	Nmgabu
N-methylcyclohexylalanine	Nmchexa	D-N-methylmethionine	Dnmmet
D-N-methylornithine	Dnmorn	N-methylcyclopentylalanine	Nmcpen
N-methylglycine	Nala	D-N-methylphenylalanine	Dnmphe
5 N-methylaminoisobutyrate	Nmaib	D-N-methylproline	Dnmpro
N-(1-methylpropyl)glycine	Nile	D-N-methylserine	Dnmser
N-(2-methylpropyl)glycine	Nleu	D-N-methylthreonine	Dnmthr
D-N-methyltryptophan	Dnmtrp	N-(1-methylethyl)glycine	Nval
D-N-methyltyrosine	Dnmtyr	N-methyla-naphylalanine	Nmanap
10 D-N-methylvaline	Dnmval	N-methylpenicillamine	Nmpen
γ -aminobutyric acid	Gabu	N-(<i>p</i> -hydroxyphenyl)glycine	Nhtyr
L- <i>t</i> -butylglycine	Tbug	N-(thiomethyl)glycine	Ncys
L-ethylglycine	Etg	penicillamine	Pen
L-homophenylalanine	Hphe	L- α -methylalanine	Mala
15 L- α -methylarginine	Marg	L- α -methylasparagine	Masn
L- α -methylaspartate	Masp	L- α -methyl- <i>t</i> -butylglycine	Mtbug
L- α -methylcysteine	Mcys	L-methylethylglycine	Metg
L- α -methylglutamine	Mgln	L- α -methylglutamate	Mglu
L- α -methylhistidine	Mhis	L- α -methylhomophenylalanine	Mhphe
20 L- α -methylisoleucine	Mile	N-(2-methylthioethyl)glycine	Nmet
L- α -methylleucine	Mleu	L- α -methyllysine	Mlys
L- α -methylmethionine	Mmet	L- α -methylnorleucine	Mnle
L- α -methylnorvaline	Mnva	L- α -methylornithine	Morn
L- α -methylphenylalanine	Mphe	L- α -methylproline	Mpro
25 L- α -methylserine	Mser	L- α -methylthreonine	Mthr
L- α -methyltryptophan	Mtrp	L- α -methyltyrosine	Mtyr
L- α -methylvaline	Mval	L-N-methylhomophenylalanine	Nmhphe

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N-(N-(2,2-diphenylethyl)	Nnbhm	N-(N-(3,3-diphenylpropyl)	Nnbhe
carbamylmethyl)glycine		carbamylmethyl)glycine	
1-carboxy-1-(2,2-diphenyl-	Nmhc		
ethylamino)cyclopropane			

5

Crosslinkers can be used, for example, to stabilise 3D conformations, using homo-bifunctional crosslinkers such as the bifunctional imido esters having $(CH_2)_n$ spacer groups with n=1 to n=6, glutaraldehyde, N-hydroxysuccinimide esters and hetero-bifunctional

10 reagents which usually contain an amino-reactive moiety such as N-hydroxysuccinimide and another group specific-reactive moiety such as maleimido or dithio moiety (SH) or carbodiimide (COOH). In addition, peptides can be conformationally constrained by, for example, incorporation of C_α and N_α-methylamino acids, introduction of double bonds between C_α and C_β atoms of amino acids and the formation of cyclic peptides or analogues

15 by introducing covalent bonds such as forming an amide bond between the N and C termini, between two side chains or between a side chain and the N or C terminus.

As stated above, these types of modifications may be important to stabilise the variant HBsAg molecule if administered to an individual or for use as a diagnostic reagent.

20

Other derivatives contemplated by the present invention include a range of glycosylation variants from a completely unglycosylated molecule to a modified glycosylated molecule. Altered glycosylation patterns may result from expression of recombinant molecules in different host cells.

25

Another aspect of the present invention extends to the variant HBsAg molecule or its recombinant, derivative or chemical form or a variant HBV comprising said HBsAg in composition form. Such compositions are particularly useful as therapeutic compositions and may be referred to herein interchangeably as biological, vaccine or pharmaceutical

30 compositions. The biological compositions are particularly useful in inducing immunological memory against infection by an HBV variant such as an HBV escape

mutant controlling by administering a variant HBsAg or a recombinant, derivative or chemical form thereof or an HBV comprising same capable of inducing an immune response including immunological memory agents.

5 Accordingly, the present invention contemplates a biological composition comprising a variant HBV or an HBsAg from said variant HBV or a recombinant or derivative form thereof or its chemical equivalent.

Generally, if an HBV is used, it is first attenuated. The biological composition according to
10 this aspect of the present invention generally further comprises one or more pharmaceutically acceptable carriers and/or diluents.

The biological composition may comprise an HBsAg or like molecule from one HBV variant or the composition may be a cocktail of HBsAgs or like molecules from a range of
15 HBV variants including the referenced HBV. Similar inclusions apply where the composition comprises an HBV.

The biological composition forms suitable for injectable use include sterile aqueous solutions (where water soluble) or sterile powders for the extemporaneous preparation of
20 sterile injectable solutions. It must be stable under the conditions of manufacture and storage and must be preserved against the contaminating action of microorganisms such as bacteria and fungi. The carrier can be a solvent or diluent containing, for example, water, ethanol, polyol (for example, glycerol, propylene glycol and liquid polyethylene glycol, and the like), suitable mixtures thereof, and vegetable oils. The preventions of the action of
25 microorganisms can be brought about by various antibacterial and antifungal agents, for example, parabens, chlorobutanol, phenol, sorbic acid, thimerosal and the like. In many cases, it will be preferable to include isotonic agents, for example, sugars or sodium chloride. Prolonged absorption of the injectable compositions can be brought about by the use in the compositions of agents delaying absorption, for example, aluminum
30 monostearate and gelatin.

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Sterile injectable solutions are prepared by incorporating the HBsAg or like molecule or HBV variant or reference strain in the required amount in the appropriate solvent or diluent as followed by sterilization such as by filter sterilization. In the case of sterile powders for the preparation of sterile injectable solutions, the preferred methods of preparation are

5 vacuum drying and the freeze-drying technique which yield a powder of the immunointeractive molecule plus any additional desired ingredient from previously sterile-filtered solution thereof. Routes of administration contemplated by the present invention including intravenous, intraperitoneal, intrathelial, subcutaneous and intracerebral.

10

The biological composition of the present invention may also be given in oral, bucal, nasal spray, inhalation, patch, drip or suppository form.

Pharmaceutically acceptable carriers and/or diluents include any and all solvents, dispersion media, coatings, antibacterial and antifungal agents, isotonic and absorption delaying agents and the like. The use of such media and agents for pharmaceutical active substances is well known in the art. Except insofar as any conventional media or agent is incompatible with the immunointeractive molecule, use thereof in the therapeutic compositions is contemplated. Supplementary active ingredients can also be incorporated into the
20 compositions.

The HBsAg or like molecule or HBV variant or reference strain will be added in a concentration effective to induce an interact immune response against the same molecule or an HBV carrying the same or an immunologically similar molecule. For example, an
25 effective amount of HBsAg may range from about 10 mg to about 2000 ng, or 50 ng to about 1000 mg or 100 ng to about 500 mg or other suitable effective amount. It is sometimes more convenient to express dosage amounts in terms of body weight. Accordingly, the effective amounts may be from, for example, about 0.5 ng/kg body weight to about 500 mg/kg body weight or an amount there between.

30

The present invention is further described by the following non-limiting Examples.

- 30 -

EXAMPLE 1
OVERLAPPING GENOME OF HBV

The overlapping genome of HBV is represented in Figure 1. The gene encoding DNA polymerase (P), overlaps the viral envelope genes, Pre-S1 and Pre-S2, and partially overlaps the X and core (C) genes. The HBV envelope comprises small, middle and large HBV surface antigens. The large protein component is referred to as the HBV surface antigen (HBsAg) and is enclosed by the S gene sequence. The Pre-S1 and Pre-S2 gene sequences encode the other envelope components.

10

EXAMPLE 2
AMINO ACID CONSENSUS SEQUENCE FO HBV DNA POLYMERASE

The amino acid consensus sequence for HBV DNA polymerase protein from genotypes A 15 through F is shown in Figure 2.

EXAMPLE 3
CONSENSUS SEQUENCE OF HBsAg

20 The nucleotide sequence from various strains of HBV encoding the surface antigen is shown in Figure 3. The amino acid sequence of the surface antigen beginning at amino acid 108 is shown above the nucleotide sequence.

EXAMPLE 4
HBsAg BINDING ASSAY

25 The effect of the Pre-S/S gene escape mutations on the binding of anti-HBs antibody is assessed using an RIA binding assay. The results are shown in Figure 4. Briefly, the expressed mutant HBsAg from transfected cell cultures is purified through a sucrose density gradient. The ability of subviral and viral particles to block the binding of wild type HBsAg to anti-HBs antibody, which does not recognise S gene escape mutants, is assessed

in an RIA format (AUSAB, Abbott). This analysis involves the binding of anti-HBs in pooled vaccine serum to increasing concentrations of wild type and mutant S protein using limiting concentrations of serum and detecting the unbound anti-HBs by AUSAB RIA.

5 The mutant S proteins analysed are shown on the right of Figure 4 together with mock and wild type HBV. As the concentration of HBsAg decreases the amount of unbound anti-HBs increase, leaving a higher anti-HBs concentration to be detected by the AUSAB assay. Even at high concentration of HBsAg from the W499Q mutant the amount of residual anti-HBs detected is similar to that of the mock transfected sample (these are represented by the
10 two curves at the top of the graph). In contrast, the amount of residual anti-HBs after binding of antibody with the other mutant HBsAg proteins is analogous to the wild type HBsAg, indicating that these variant vHBsAg proteins recognise the anti-HBs with similar efficiency as the wild type protein.

15 Two of the mutant S proteins (Figure 4: V519L and the triple mutant which contains the mutations V519L + L526M +M550V with respect to the polymerase protein in the overlapping reading frame) had partial binding of anti-HBsAg. The binding efficacy of the mutant S proteins to HBsAg is altered when compared to wild type HBsAg. This suggests that viruses carrying these mutations may not be detected by anti-HBsAg as efficiently as
20 wild type virus and thus may escape immune detection. Hepatitis B virus with these and/or other HBsAg mutations, which have partial binding to anti-HBsAg, may also escape immune detection and protection.

The dual mutant in Figure 4 represents L526M/M550V while the triple mutant represents
25 V519L/L526M/M550V.

EXAMPLE 5

HBV VARIANTS PRODUCED BY SITE DIRECTED MUTAGENESIS

30 Table 2 provides a summary of some of the HBV variants produced by site directed mutagenesis.

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EXAMPLE 6
FCV MUTATION

Table 3 provides a summary of mutations induced by famciclovir (FCV).

5

Those skilled in the art will appreciate that the invention described herein is susceptible to variations and modifications other than those specifically described. It is to be understood that the invention includes all such variations and modifications. The invention also includes all of the steps, features, compositions and compounds referred to
10 or indicated in this specification, individually or collectively, and any and all combinations of any two or more of said steps or features.

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TABLE 2
HBV VARIANTS PRODUCED BY SITE DIRECTED MUTAGENESIS

Nucleotide analogue selected

Polymerase mutations

	Corresponding surface S mutation
1. G499E (B domain)	D144E
2. W499Q (B domain)	G145R
3. F512L (B domain)	A157D
4. V519L (B domain)	E164D
5. L526M (B domain)	no change
6. M550V (C domain)	M195I
7. M550V (C domain)	W196L
8. M550I (C domain)	S196W
9. V553I (C domain)	M198I
10. V553I (C domain)	W199S
11. S565P	S210R

Double polymerase mutation

	Corresponding S mutation
12. L526M/M550V	M195I
13. L526M/M550I	S196W

Triple polymerase mutation

	Corresponding S mutation
14. V519L/L526M/M550V	E164D, M195I, W196L
15. VS19L/L526M/M550I	E164D, S196W

HBsAg escape mutant

	Corresponding HBV polymerase changes
16. K122R (loop 1 "a" determinant)	Q476P
17. T126S (loop 1 "a" determinant)	N480G
18. T131N (loop 1 "a" determinant)	N485K
19. K141E (loop 2 "a" determinant)	K495R
20. G145K (loop 2 "a" determinant)	R499Q
21. R160N	I515L

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TABLE 3

FCV Mutations		Number of patients with Mutation (%)	
S421L	A Domain	1/34	3%
N422K	A Domain	1/34	3%
L423L/M/V	A Domain	1/34	3%
S424T	A Domain	2/34	6%
S/D 455P		7/34*	20.5%
N464D		1/34	3%
Q471K/N		2/34	6%
D/N 480E		1/34	3%
T484H		1/34	3%
R499K		4/34	12%
V519L	B Domain	3/34	9%
L/M/V523L	B Domain	1/34	3%
F524L/F	B Domain	1/34	3%
L526M	B Domain	5/34	15%
A527T	B Domain	1/34	3%
I533I/V		2/34	6%
V537I		1/34	3%
S565A		1/34	3%
S/D576F/S	D Domain	1/34	3%
L593V		1/34	3%
H/Y594H	E Domain	1/34	3%
T/M596M	E Domain	1/34	3%

* Only detected in BMT patients on FCV

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CLAIMS

1. A variant Hepatitis B Virus (HBV) comprising a surface component exhibiting an altered immunological profile compared to a reference HBV.
2. A variant HBV according to claim 1 wherein the surface component on the variant is a surface antigen which comprises a single or multiple amino acid substitution, addition and/or deletion or truncation compared to a surface antigen from said reference HBV and wherein an antibody generated to the reference surface antigen from the reference HBV exhibits reduced capacity for neutralizing said variant HBV.
3. A variant HBV according to claim 2 wherein the surface antigen on the variant comprises an amino acid substitution, addition and/or deletion or truncation compared to the amino acid sequence setforth below:

M X₁ X₂ X₃ X₄ S G X₅ L X₆ P L X₇ V L Q A X₈ X₉ F X₁₀ L T X₁₁ I X₁₂ X₁₃ I P
X₁₄ S L X₁₅ S W W T S L N F L G X₁₆ X₁₇ X₁₈ X₁₉ C X₂₀ G X₂₁ N X₂₂ Q S
X₂₃ X₂₄ S X₂₅ H X₂₆ P X₂₇ X₂₈ C P P X₂₉ C X₃₀ G Y R W M C L X₃₁ R F I I
F L X₃₂ I L L L C L I F L L V L L D X₃₃ Q G M L X₃₄ V C P L X₃₅ P X₃₆ X₃₇
X₃₈ T T S X₃₉ X₄₀ X₄₁ C X₄₂ T C X₄₃ X₄₄ X₄₅ X₄₆ Q G X₄₇ S X₄₈ X₄₉ P X₅₀ X₅₁
C C X₅₂ K P X₅₃ X₅₄ G N C T C I P I P S X₅₅ W A X₅₆ X₅₇ X₅₈ X₅₉ L W E
X₆₀ X₆₁ S X₆₂ R X₆₃ S W L X₆₄ L L X₆₅ X₆₆ F V Q X₆₇ X₆₈ X₆₉ X₇₀ L X₇₁ P X₇₂ V
W X₇₃ X₇₄ X₇₅ I W X₇₆ X₇₇ W X₇₈ W X₇₉ P X₈₀ X₈₁ X₈₂ X₈₃ I X₈₄ X₈₅ P F X₈₆
P L L P I F X₈₇ X₈₈ L X₈₉ X₉₀ X₉₁ I [Formula I];

wherein:

X₁ is E or G or D;

X₂ is N or S or K;

X₃ is I or T;

X₄ is T or A;
X₅ is F or L;
X₆ is G or R;
X₇ is L or R;
X₈ is G or V;
X₉ is F or C;
X₁₀ is L or S or W;
X₁₁ is R or K;
X₁₂ is L or R;
X₁₃ is T or K;
X₁₄ is Q or K;
X₁₅ is D or H;
X₁₆ is G or E or A;
X₁₇ is S or A or V or T or L;
X₁₈ is P or T;
X₁₉ is V or R or T or K or G;
X₂₀ is L or P;
X₂₁ is Q or L or K;
X₂₂ is S or L;
X₂₃ is P or Q;
X₂₄ is T or I;
X₂₅ is N or S;
X₂₆ is S or L;
X₂₇ is T or I;
X₂₈ is S or C;
X₂₉ is I or T;
X₃₀ is P or A;

X_{31} is R or Q;
 X_{32} is F or C;
 X_{33} is Y or C;
 X_{34} is P or H or S;
 X_{35} is I or L;
 X_{36} is G or R;
 X_{37} is S or T;
 X_{38} is T or S;
 X_{39} is T or V or A;
 X_{40} is G or E or Q;
 X_{41} is P or A or S;
 X_{42} is K or R;
 X_{43} is T or M;
 X_{44} is T or I or S or A;
 X_{45} is P or T or A or I or L;
 X_{46} is A or V;
 X_{47} is N or T;
 X_{48} is M or K or L;
 X_{49} is F or Y or I;
 X_{50} is S or Y;
 X_{51} is C or S;
 X_{52} is T or I or S;
 X_{53} is T or S;
 X_{54} is D or A;
 X_{55} is S or T;
 X_{56} is F or L;
 X_{57} is A or G or V;

X₅₈ is K or R or T;
X₅₉ is Y or F;
X₆₀ is W or G;
X₆₁ is A or G;
X₆₂ is V or A;
X₆₃ is F or L;
X₆₄ is S or N;
X₆₅ is V or A;
X₆₆ is P or Q;
X₆₇ is W or C or S;
X₆₈ is F or C;
X₆₉ is V or D or A;
X₇₀ is G or E;
X₇₁ is S or F;
X₇₂ is T or I;
X₇₃ is L or P;
X₇₄ is S or L;
X₇₅ is A or V;
X₇₆ is M or I;
X₇₇ is M or I;
X₇₈ is Y or F;
X₇₉ is G or E;
X₈₀ is S or N or K;
X₈₁ is L or Q;
X₈₂ is Y or F or H or C;
X₈₃ is S or G or N or D or T;
X₈₄ is V or L;

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X_{85} is S or N;
 X_{86} is I or M or L;
 X_{87} is F or C;
 X_{88} is C or Y;
 X_{89} is W or R;
 X_{90} is V or A; and
 X_{91} is Y or I or S;

and wherein the variant HBV is selected for by a nucleotide analogue of HBV DNA polymerase.

4. A variant HBV according to claim 2 wherein the surface antigen on the variant comprises an amino acid substitution, addition and/or deletion or truncation compared to the amino acid sequence setforth below:

M X_1 X_2 X_3 X_4 S G X_5 L X_6 P L X_7 V L Q A X_8 X₉ F X_{10} L T X_{11} I X_{12} X₁₃ I P
 X_{14} S L X_{15} S W W T S L N F L G X_{16} X₁₇ X₁₈ X₁₉ C X_{20} G X_{21} N X_{22} Q S
 X_{23} X₂₄ S X_{25} H X_{26} P X_{27} X₂₈ C P P X_{29} C X_{30} G Y R W M C L X_{31} R F I I
F L X_{32} I L L L C L I F L L V L L D X_{33} Q G M L X_{34} V C P L X_{35} P X_{36} X₃₇
 X_{38} T T S X_{39} X₄₀ C X_{41} C X_{42} T C X_{43} X₄₄ X₄₅ X₄₆ Q G X_{47} S X_{48} X₄₉ P X_{50} X₅₁
C C X_{52} K P X_{53} X₅₄ G N C T C I P I P S X_{55} W A X_{56} X₅₇ X₅₈ X₅₉ L W E
 X_{60} X₆₁ S X_{62} R X_{63} S W L X_{64} LLX₆₅ X_{66} F V Q X_{67} X₆₈ X₆₉ X₇₀ L X_{71} P X_{72} V
W
 X_{73} X₇₄ X₇₅ I W X_{76} X₇₇ W X_{78} W X_{79} P X_{80} X₈₁ X₈₂ X₈₃ I X_{84} X₈₅ P F X_{86} P L
L P I F X_{87} X₈₈ L X_{89} X₉₀ X₉₁ I [Formula I];

wherein:

X_1 is E or G or D;
 X_2 is N or S or K;
 X_3 is I or T;
 X_4 is T or A;
 X_5 is F or L;
 X_6 is G or R;
 X_7 is L or R;
 X_8 is G or V;
 X_9 is F or C;
 X_{10} is L or S or W;
 X_{11} is R or K;
 X_{12} is L or R;
 X_{13} is T or K;
 X_{14} is Q or K;
 X_{15} is D or H;
 X_{16} is G or E or A;
 X_{17} is S or A or V or T or L;
 X_{18} is P or T;
 X_{19} is V or R or T or K or G;
 X_{20} is L or P;
 X_{21} is Q or L or K;
 X_{22} is S or L;
 X_{23} is P or Q;
 X_{24} is T or I;
 X_{25} is N or S;
 X_{26} is S or L;

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X₂₇ is T or I;
X₂₈ is S or C;
X₂₉ is I or T;
X₃₀ is P or A;
X₃₁ is R or Q;
X₃₂ is F or C;
X₃₃ is Y or C;
X₃₄ is P or H or S;
X₃₅ is I or L;
X₃₆ is G or R;
X₃₇ is S or T;
X₃₈ is T or S;
X₃₉ is T or V or A;
X₄₀ is G or E or Q;
X₄₁ is P or A or S;
X₄₂ is K or R;
X₄₃ is T or M;
X₄₄ is T or I or S or A;
X₄₅ is P or T or A or I or L;
X₄₆ is A or V;
X₄₇ is N or T;
X₄₈ is M or K or L;
X₄₉ is F or Y or I;
X₅₀ is S or Y;
X₅₁ is C or S;
X₅₂ is T or I or S;
X₅₃ is T or S;

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X_{54} is D or A;
 X_{55} is S or T;
 X_{56} is F or L;
 X_{57} is A or G or V;
 X_{58} is K or R or T;
 X_{59} is Y or F;
 X_{60} is W or G;
 X_{61} is A or G;
 X_{62} is V or A;
 X_{63} is F or L;
 X_{64} is S or N;
 X_{65} is V or A;
 X_{66} is P or Q;
 X_{67} is W or C or S;
 X_{68} is F or C;
 X_{69} is V or D or A;
 X_{70} is G or E;
 X_{71} is S or F;
 X_{72} is T or I;
 X_{73} is L or P;
 X_{74} is S or L;
 X_{75} is A or V;
 X_{76} is M or I;
 X_{77} is M or I;
 X_{78} is Y or F;
 X_{79} is G or E;
 X_{80} is S or N or K;

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X₈₁ is L or Q;
X₈₂ is Y or F or H or C;
X₈₃ is S or G or N or D or T;
X₈₄ is V or L;
X₈₅ is S or N;
X₈₆ is I or M or L;
X₈₇ is F or C;
X₈₈ is C or Y;
X₈₉ is W or R;
X₉₀ is V or A; and
X₉₁ is Y or I or S.

and wherein the variant HBV is selected for following immunological therapy directed against a surface antigen as defined in Formula I.

5. A variant HBV according to claim 1 comprising a nucleotide sequence having a single or multiple nucleotide substitution, addition and/or deletion or truncation of the nucleotide sequence as set forth in formula III below:

AC N₁ A A A C C T N₂ N₃ G G A N₄ G G A A A N₅ T G C A C N₆ T G T
A T T C C C A T C C C A T C N₇ T C N₈ T G G G C T T T C G N₉ A A
N₁₀ A T N₁₁ C C T A T G G G A G N₁₂ G G G C C T C A G N₁₃ C C G T
T T C T C N₁₄ T G G C T C A G T T A C T A G T G C C A T T T G T
T C A G T G G T T C G N₁₅ A G G G C T T T C C C C A C T G T N₁₆
T G G C T T T C A G N₁₇ T A T A T G G A T G A T G T G G T N₁₈ T T
G G G G G C C A A G T C T G T A C A N₁₉ C A T C N₂₀ T G A G T C C
C T T T N₂₁ T N₂₂ C C N₂₃ C T N₂₄ T T A C C A A T T T C T T N₂₅ T G
T C T N₂₆ T G G G N₂₇ A T A C A T T [Formula III];

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wherein:

N₁ is A or C;
N₂ is T or A;
N₃ is C or T;
N₄ is C or T;
N₅ is C or T;
N₆ is C or T;
N₇ is A or G;
N₈ is T or C;
N₉ is C or G;
N₁₀ is G or A;
N₁₁ is T or A;
N₁₂ is T or G;
N₁₃ is T or C;
N₁₄ is C or T;
N₁₅ is T or C;
N₁₆ is T or C;
N₁₇ is T or C;
N₁₈ is A or T;
N₁₉ is A or G;
N₂₀ is T or G;
N₂₁ is A or T;
N₂₂ is A or G;
N₂₃ is T or G;
N₂₄ is A or G;
N₂₅ is T or C;
N₂₆ is T or C; and

N_{27} is T or C.

and wherein the HBV variant has a surface antigen exhibiting an altered immunological profile relative to the surface antigen as defined in Formula I.

6. An isolated variant Hepatitis B virus surface antigen (HBsAg) or a recombinant form thereof or a derivative or chemical equivalent thereof wherein said HBsAg or its recombinant or derivative form or its chemical equivalent exhibits an altered immunological profile compared to an HBsAg from a reference HBV.

7. An isolated variant HBsAg or a recombinant or derivative form thereof or a chemical equivalent thereof according to claim 6 wherein the variant HBsAg comprises an amino acid sequence having a single or multiple substitution, addition and/or deletion or truncation of the amino acid sequence set forth below:

M X₁ X₂ X₃ X₄ S G X₅ L X₆ P L X₇ V L Q A X₈ X₉ F X₁₀ L T X₁₁ I X₁₂ X₁₃ I P
X₁₄ S L X₁₅ S W W T S L N F L G X₁₆ X₁₇ X₁₈ X₁₉ C X₂₀ G X₂₁ N X₂₂ Q S
X₂₃ X₂₄ S X₂₅ H X₂₆ P X₂₇ X₂₈ C P P X₂₉ C X₃₀ G Y R W M C L X₃₁ R F I I
F L X₃₂ I L L L C L I F L L V L L D X₃₃ Q G M L X₃₄ V C P L X₃₅ P X₃₆ X₃₇
X₃₈ T T S X₃₉ X₄₀ X₄₁ C X₄₂ T C X₄₃ X₄₄ X₄₅ X₄₆ Q G X₄₇ S X₄₈ X₄₉ P X₅₀ X₅₁
C C X₅₂ K P X₅₃ X₅₄ G N C T C I P I P S X₅₅ W A X₅₆ X₅₇ X₅₈ X₅₉ L W E
X₆₀ X₆₁ S X₆₂ R X₆₃ S W L X₆₄ LLX₆₅ X₆₆ F V Q X₆₇ X₆₈ X₆₉ X₇₀ L X₇₁ P X₇₂ V
W
X₇₃ X₇₄ X₇₅ I W X₇₆ X₇₇ W X₇₈ W X₇₉ P X₈₀ X₈₁ X₈₂ X₈₃ I X₈₄ X₈₅ P F X₈₆ P L
L P I F X₈₇ X₈₈ L X₈₉ X₉₀ X₉₁ I [Formula I];

wherein:

X₁ is E or G or D;

X₂ is N or S or K;

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X₃ is I or T;
X₄ is T or A;
X₅ is F or L;
X₆ is G or R;
X₇ is L or R;
X₈ is G or V;
X₉ is F or C;
X₁₀ is L or S or W;
X₁₁ is R or K;
X₁₂ is L or R;
X₁₃ is T or K;
X₁₄ is Q or K;
X₁₅ is D or H;
X₁₆ is G or E or A;
X₁₇ is S or A or V or T or L;
X₁₈ is P or T;
X₁₉ is V or R or T or K or G;
X₂₀ is L or P;
X₂₁ is Q or L or K;
X₂₂ is S or L;
X₂₃ is P or Q;
X₂₄ is T or I;
X₂₅ is N or S;
X₂₆ is S or L;
X₂₇ is T or I;
X₂₈ is S or C;
X₂₉ is I or T;

X_{30} is P or A;
 X_{31} is R or Q;
 X_{32} is F or C;
 X_{33} is Y or C;
 X_{34} is P or H or S;
 X_{35} is I or L;
 X_{36} is G or R;
 X_{37} is S or T;
 X_{38} is T or S;
 X_{39} is T or V or A;
 X_{40} is G or E or Q;
 X_{41} is P or A or S;
 X_{42} is K or R;
 X_{43} is T or M;
 X_{44} is T or I or S or A;
 X_{45} is P or T or A or I or L;
 X_{46} is A or V;
 X_{47} is N or T;
 X_{48} is M or K or L;
 X_{49} is F or Y or I;
 X_{50} is S or Y;
 X_{51} is C or S;
 X_{52} is T or I or S;
 X_{53} is T or S;
 X_{54} is D or A;
 X_{55} is S or T;
 X_{56} is F or L;

X₅₇ is A or G or V;
X₅₈ is K or R or T;
X₅₉ is Y or F;
X₆₀ is W or G;
X₆₁ is A or G;
X₆₂ is V or A;
X₆₃ is F or L;
X₆₄ is S or N;
X₆₅ is V or A;
X₆₆ is P or Q;
X₆₇ is W or C or S;
X₆₈ is F or C;
X₆₉ is V or D or A;
X₇₀ is G or E;
X₇₁ is S or F;
X₇₂ is T or I;
X₇₃ is L or P;
X₇₄ is S or L;
X₇₅ is A or V;
X₇₆ is M or I;
X₇₇ is M or I;
X₇₈ is Y or F;
X₇₉ is G or E;
X₈₀ is S or N or K;
X₈₁ is L or Q;
X₈₂ is Y or F or H or C;
X₈₃ is S or G or N or D or T;

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X_{84} is V or L;
 X_{85} is S or N;
 X_{86} is I or M or L;
 X_{87} is F or C;
 X_{88} is C or Y;
 X_{89} is W or R;
 X_{90} is V or A; and
 X_{91} is Y or I or S;

and wherein a neutralizing antibody directed to a reference HBV exhibits no or reduced neutralizing activity to an HBV carrying said variant HBsAg.

8. An isolated variant HBsAg according to claim 7 encoded by a nucleotide sequence having a single or multiple nucleotide substitution, addition and/or deletion or truncation relative to the nucleotide sequence set forth below:

A C N₁ A A A C C T N₂ N₃ G G A N₄ G G A A A N₅ T G C A C N₆ T G T A
T T C C C A T C C C A T C N₇ T C N₈ T G G G C T T T C G N₉ A A N₁₀ A
T N₁₁ C C T A T G G G A G N₁₂ G G G C C T C A G N₁₃ C C G T T T C T C
N₁₄ T G G C T C A G T T A C T A G T G C C A T T T G T T C A G T G G
T T C G N₁₅ A G G G C T T C C C C A C T G T N₁₆ T G G C T T T C A
G N₁₇ T A T A T G G A T G A T G T G G T N₁₈ T T G G G G G C C A A G
T C T G T A C A N₁₉ C A T C N₂₀ T G A G T C C C T T T N₂₁ T N₂₂ C C N₂₃
C T N₂₄ T T A C C A A T T T C T T N₂₅ T G T C T N₂₆ T G G G N₂₇ A T A
C A T T [FORMULA III];

wherein:

N₁ is A or C;

N₂ is T or A;
N₃ is C or T;
N₄ is C or T;
N₅ is C or T;
N₆ is C or T;
N₇ is A or G;
N₈ is T or C;
N₉ is C or G;
N₁₀ is G or A;
N₁₁ is T or A;
N₁₂ is T or G;
N₁₃ is T or C;
N₁₄ is C or T;
N₁₅ is T or C;
N₁₆ is T or C;
N₁₇ is T or C;
N₁₈ is A or T;
N₁₉ is A or G;
N₂₀ is T or G;
N₂₁ is A or T;
N₂₂ is A or G;
N₂₃ is T or G;
N₂₄ is A or G;
N₂₅ is T or C;
N₂₆ is T or C; and
N₂₇ is T or C.

9. A variant HBV or an isolated HBsAg from said variant HBV wherein said variant HBV comprises an HBsAg having a single or multiple amino acid substitution, addition and/or deletion or truncation relative to the HBsAg on a reference HBV and whereas the HBsAg variant is defined by a single or multiple amino acid substitution, addition and/or truncation to the catalytic region of HBV DNA polymerase as defined below:

S Z₁ L S W L S L D V S A A F Y H Z₂ P L H P A A M P H L L Z₃ G S S
G L Z₄ R Y V A R L S S Z₅ S Z₆ Z₇ X N Z₈ Q Z₉ Z₁₀ X X X Z₁₁ L H Z₁₂ Z₁₃
C S R Z₁₄ L Y V S L Z₁₅ L L Y Z₁₆ T Z₁₇ G Z₁₈ K L H L Z₁₉ Z₂₀ H P I Z₂₁ L
G F R K Z₂₂ P M G Z₂₃ G L S P F L L A Q F T S A I Z₂₄ Z₂₅ Z₂₆ Z₂₇ Z₂₈ R
A F Z₂₉ H C Z₃₀ Z₃₁ F Z₃₂ Y M * D D Z₃₃ V L G A Z₃₄ Z₃₅ Z₃₆ Z₃₇ H Z₃₈ E Z₃₉
L Z₄₀ Z₄₁ Z₄₂ Z₄₃ Z₄₄ Z₄₅ Z₄₆ L L Z₄₇ Z₄₈ G I H L N P Z₄₉ K T K R W G Y S
L N F M G Y Z₅₀ I G [Formula II];

wherein:

X is any amino acid;

Z₁ is N or D;

Z₂ is I or P;

Z₃ is I or V;

Z₄ is S or D;

Z₅ is T or N;

Z₆ is R or N;

Z₇ is N or I;

Z₈ is N or Y or H;

Z₉ is H or Y;

Z₁₀ is G or R;

Z₁₁ is D or N;

Z₁₂ is D or N;

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Z_{13} is S or Y;
 Z_{14} is N or Q;
 Z_{15} is L or M;
 Z_{16} is K or Q;
 Z_{17} is Y or F;
 Z_{18} is R or W;
 Z_{19} is Y or L;
 Z_{20} is S or A;
 Z_{21} is I or V;
 Z_{22} is I or L;
 Z_{23} is V or G;
 Z_{24} is C or L;
 Z_{25} is A or S;
 Z_{26} is V or M;
 Z_{27} is V or T;
 Z_{28} is R or C;
 Z_{29} is F or P;
 Z_{30} is L or V;
 Z_{31} is A or V;
 Z_{32} is S or A;
 Z_{33} is V or L or M;
 Z_{34} is K or R;
 Z_{35} is S or T;
 Z_{36} is V or G;
 Z_{37} is Q or E;
 Z_{38} is L or S or R;
 Z_{39} is S or F;

Z_{40} is F or Y;
 Z_{41} is T or A;
 Z_{42} is A or S;
 Z_{43} is V or I;
 Z_{44} is T or C;
 Z_{45} is N or S;
 Z_{46} is F or V;
 Z_{47} is S or D;
 Z_{48} is L or V;
 Z_{49} is N or Q;
 Z_{50} is V or I; and
 M^* is amino acid 550.

10. A variant HBV or variant HBsAg from said variant HBV comprising a mutation selected from the list consisting of G112R, T123P, Y/F134S, D144E, G145R, A157D, E164D, F170L, M195I, W196L, S196W, W196STOP, M198I, W199S, S204T and S210R wherein "STOP" means a stop codon.

11. A variant HBV or variant HBsAg from said variant HBV comprising a mutation selected from the list consisting of :
D144E, G145R, A157D, E164D, M195I, W196L, S196W, W196STOP, M198I, W199S and S210R wherein "STOP" means a codon.

12. A variant HBV or variant HBsAg from said variant HBV comprising a mutation selected from the list consisting of:
Q476, N480G, N485K, K495R, R499O, G499E, W499Q, F512L, I515L, V519L, L526M, M550V, M550I, V553I and S565P.

13. A composition comprising a variant HBV or variant HBsAg according to any one of

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claims 1 to 12 or a recombinant or derivative form or its chemical equivalent.

14. A composition according to claim 13 further comprising one or more pharmaceutically acceptable carriers and/or diluents.

15. A method for the treatment or prophylaxis of HBV infection said method comprising administering to a subject an amount of a variant HBV or variant HBsAg according to any one of claims 1 to 12 or a composition according to claim 13 or 14, said amount being effective to induce an immune response to said variant HBV.

16. Use of a variant HBV or an HBsAg from said HBV in the manufacture of a medicament for the treatment or prophylaxis of infection by said variant HBV.

17. Use of a variant HBV or an HBsAg from said variant HBV in screening for an agent useful in the treatment or prophylaxis of infection by said variant HBV.

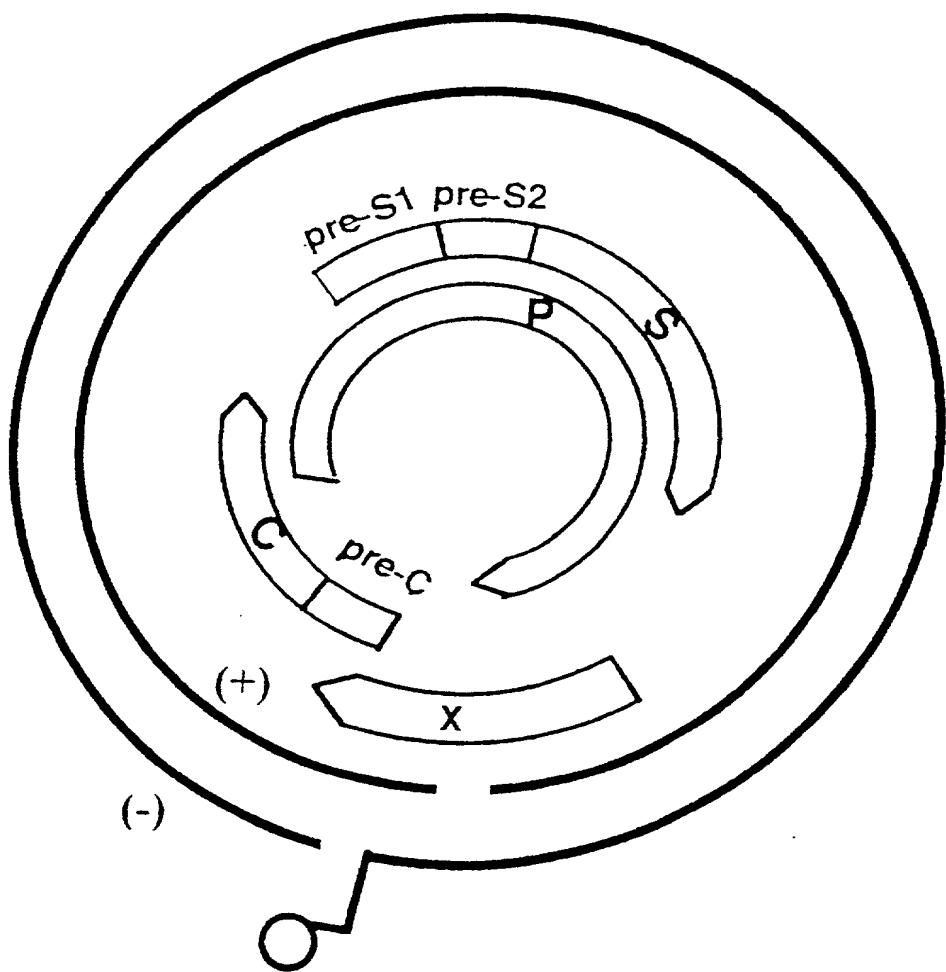


Figure 1
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PCT/AU99/00993

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(421) 430 440 450

422

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SNDLSWLSDL VSAAFYH^I_PPL HPAAMPPLLIV GSSGLD_SRYVA
Domain A

HBSAg_G112R T123P Y/F134S D144E_G145R
460 470 480 490
464 466 477 488 499
RLSST_NSR_{NNI}*N NYH_HQY_HG_R***D_NLH D_NY_SCSR_D_QLYVS LL_MLLY_K_QTY_FGR_W

HBSAg A157D E164D F170L
500 510 520 530
512 519 523/524/526/528/530
KLHL_Y_LS_AHPI_I_V LGFRKI_LPMGV_G GLSPFLLAQF TSAIC_LS_AV_MV_TR_CR
Domain B

W196L W199S
HBSAg M195I/S196W M198T S204T S210R
540 550 560
546 550 553 559 565
AFF_PHCL_V_AvFS_AY MDDV_L_MV_LGAK_R_S_T V_G_QEHL_S_RES_FLY_FTA_S_A
Domain C

570 580 590

575

I_V_T_C_N_S_F_VLL_S_D_L_VGI HLNPN_QKTKRW GYSLNFMGY_I_V_I G
Domain D Domain E

Figure 2

Substitute Sheet (Rule 26) RO/AU

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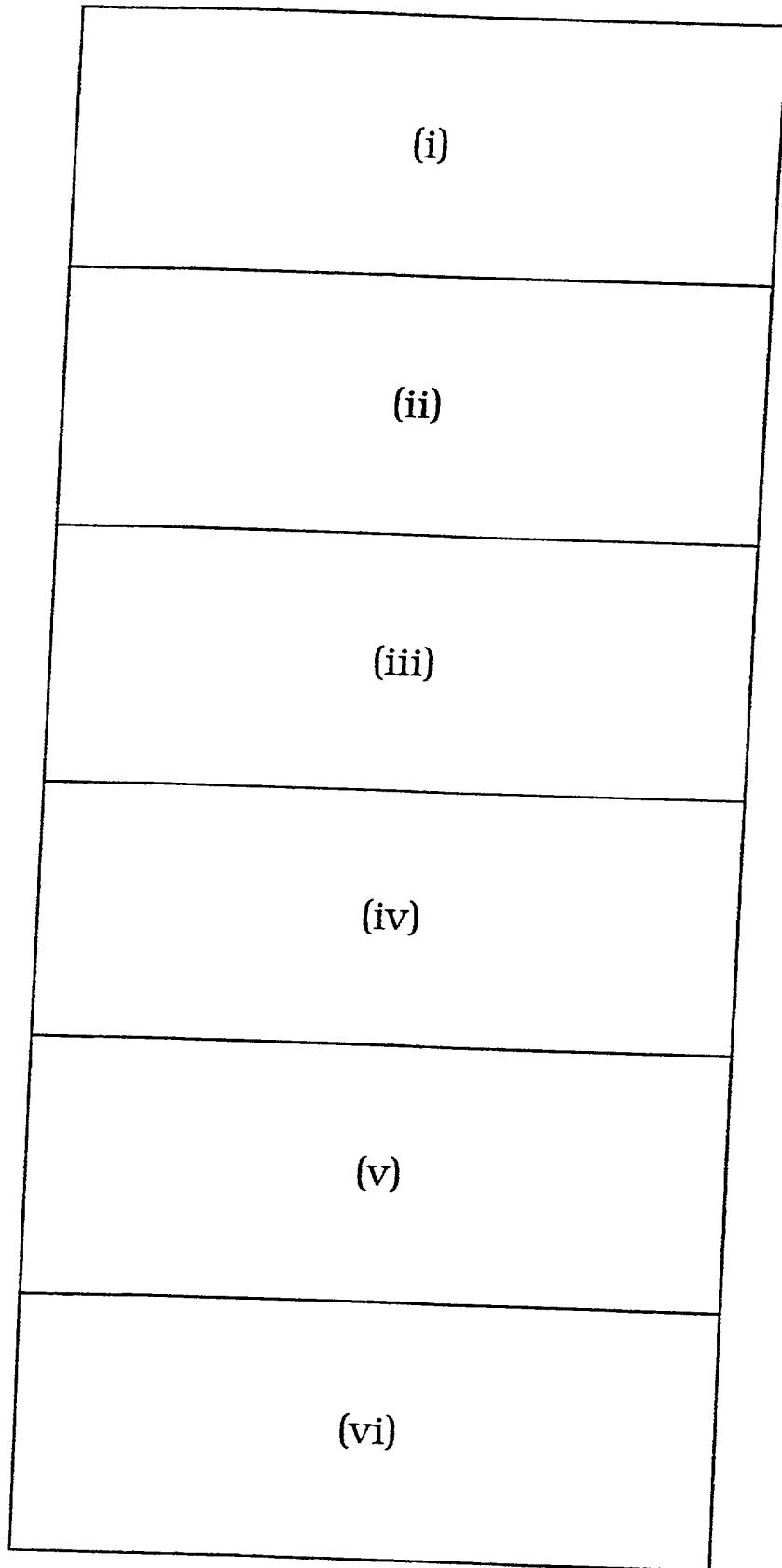


Figure 3
Substitute Sheet (Rule 26) RO/AU

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PCT/AU99/00993

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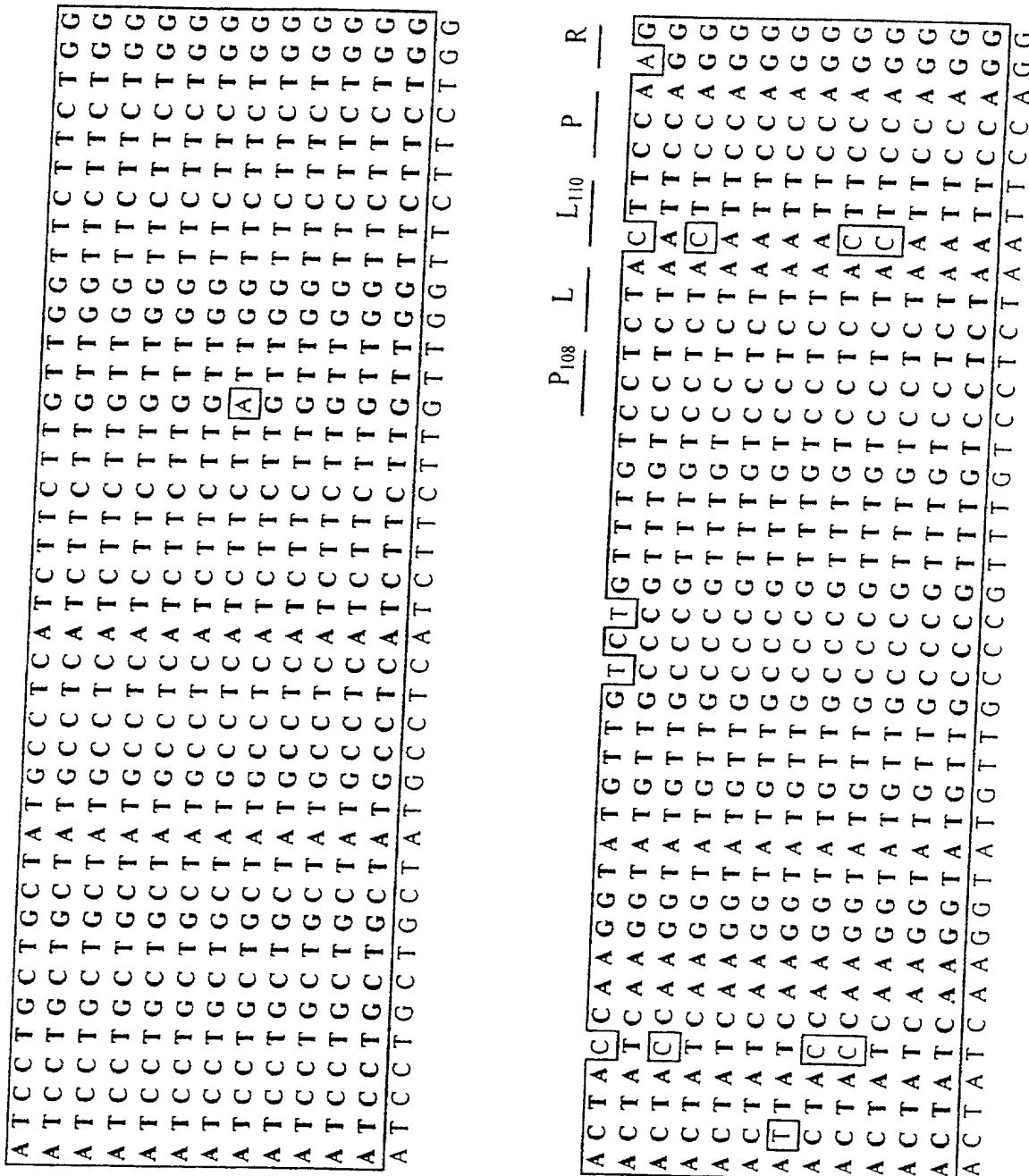


Figure 3(j)

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*329616/HPBADR1CG
221499/HPBADW3
221500/HPBCG
62280/XXHEPAV
59439/HBVAYWE
59429/HBVAYWC
59418/HBVADW2
59408/HBVADRM
59404/HBVADR4
3329640/HPBAYW
313780/HBVAYMMCG
22229417/HPBADW1

Substitute Sheet (Rule 26) RO/AU

Figure 3(ii)

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*329616/HPBADRCG
2221499/HPBADW3
2221500/HPBCG
62280/XXHEPAV
59439/HBVAYWE
59429/HBVAYWC
59418/HBVADW2
59408/HBVADM
59404/HBVADR4
3229640/HPBAYW
313780/HBVAYWMC
22229417/HPBADW1

Figure 3(iii)

09/831686

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PCT/AU99/00993

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*329616/HPBADR1CG	*329616/HPBADR1CG
221499/HPBADW3	221499/HPBADW3
221500/HPBCG	221500/HPBCG
62280/XXHEPAV	62280/XXHEPAV
59439/HBVAYWE	59439/HBVAYWE
59429/HBVAYWC	59429/HBVAYWC
59418/HBVADW2	59418/HBVADW2
59408/HBVADRM	59408/HBVADRM
59404/HBVADR4	59404/HBVADR4
329640/HPBAYW	329640/HPBAYW
313780/HBVAYWMCG	313780/HBVAYWMCG
229417/HPBADW1	229417/HPBADW1
*329616/HPBADR1CG	*329616/HPBADR1CG
221499/HPBADW3	221499/HPBADW3
221500/HPBCG	221500/HPBCG
62280/XXHEPAV	62280/XXHEPAV
59439/HBVAYWE	59439/HBVAYWE
59429/HBVAYWC	59429/HBVAYWC
59418/HBVADW2	59418/HBVADW2
59408/HBVADRM	59408/HBVADRM
59404/HBVADR4	59404/HBVADR4
329640/HPBAYW	329640/HPBAYW
313780/HBVAYWMCG	313780/HBVAYWMCG
229417/HPBADW1	229417/HPBADW1

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S ₁₉₃	V ₁₉₄	I ₁₉₅	W ₁₉₆	M ₁₉₇	W ₁₉₉	Y ₂₂₀	W	G	P	S	L	
T T C A G T T A T G G A T G A T G G T A T T G G G C C A A G T C T G	T T C A G T T A T G G A T G A T G G T A T T G G G C C A A G T C T G	T T C A G T T A T G G A T G A T G G T A T T G G G C C A A G T C T G	T T C A G T T A T G G A T G A T G G T A T T G G G C C A A G T C T G	T T C A G T T A T G G A T G A T G G T A T T G G G C C A A G T C T G	T T C A G T T A T G G A T G A T G G T A T T G G G C C A A G T C T G	T T C A G T T A T G G A T G A T G G T A T T G G G C C A A G T C T G	T T C A G T T A T G G A T G A T G G T A T T G G G C C A A G T C T G	T T C A G T T A T G G A T G A T G G T A T T G G G C C A A G T C T G	T T C A G T T A T G G A T G A T G G T A T T G G G C C A A G T C T G	T T C A G T T A T G G A T G A T G G T A T T G G G C C A A G T C T G	T T C A G T T A T G G A T G A T G G T A T T G G G C C A A G T C T G	
59404/HBVADRM												
329616/HPBADR1CG	221499/HPBADW3	221500/HPBCG	622280/XXHEPAV	59439/HBVAYWE	59429/HBVAYWC	59418/HBVADW2	59408/HBVADRM	329640/HPBAYW	313780/HBVAYWMCG	229417/HPBADW1		
T A C A A C A T C T T G A G T C C C T T A C C T C T A T T A C C A A T T T	T A C A A C A T C T T G A G T C C C T T A C C T C T A T T A C C A A T T T	T A C A A C A T C T T G A G T C C C T T A C C T C T A T T A C C A A T T T	T A C A A C A T C T T G A G T C C C T T A C C T C T A T T A C C A A T T T	T A C A A C A T C T T G A G T C C C T T A C C T C T A T T A C C A A T T T	T A C A A C A T C T T G A G T C C C T T A C C T C T A T T A C C A A T T T	T A C A A C A T C T T G A G T C C C T T A C C T C T A T T A C C A A T T T	T A C A A C A T C T T G A G T C C C T T A C C T C T A T T A C C A A T T T	T A C A A C A T C T T G A G T C C C T T A C C T C T A T T A C C A A T T T	T A C A A C A T C T T G A G T C C C T T A C C T C T A T T A C C A A T T T	T A C A A C A T C T T G A G T C C C T T A C C T C T A T T A C C A A T T T	T A C A A C A T C T T G A G T C C C T T A C C T C T A T T A C C A A T T T	
622280/XXHEPAV	59439/HBVAYWE	59429/HBVAYWC	59418/HBVADW2	59408/HBVADRM	59404/HBVADRM	329640/HPBAYW	313780/HBVAYWMCG	229417/HPBADW1				

*329616/HPBADR1CG
221499/HPBADW3
221500/HPBCG
622280/XXHEPAV
59439/HBVAYWE
59429/HBVAYWC
59418/HBVADW2
59408/HBVADRM
59404/HBVADRM
329640/HPBAYW
313780/HBVAYWMCG
229417/HPBADW1

Figure 3(v)

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Figure 3(vi)

Substitute Sheet (Rule 26) RO/AU

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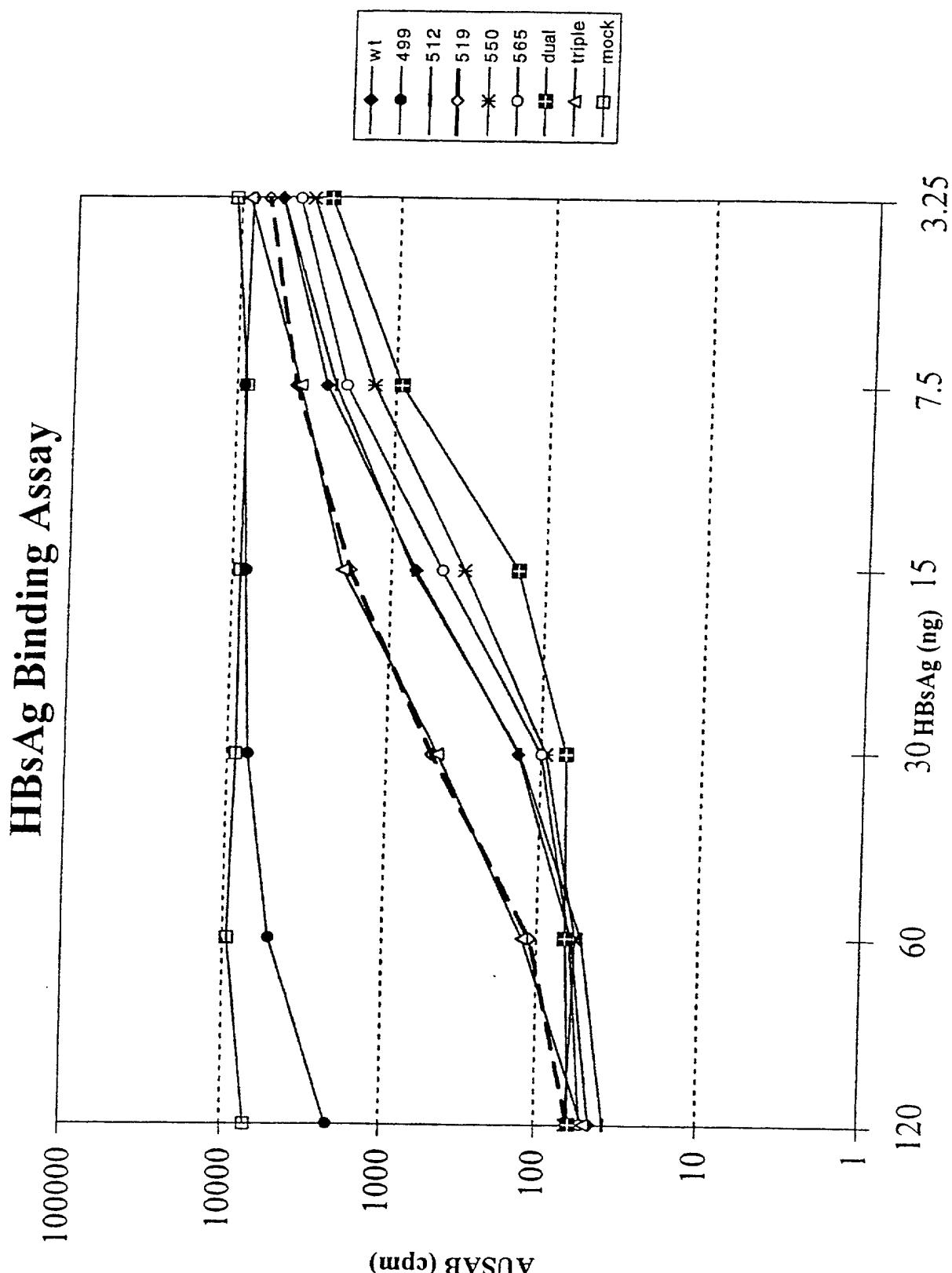


Figure 4

PATENT
Attorney Docket No. 37921-151956

DECLARATION AND POWER OF ATTORNEY

As a below named inventor, I hereby declare that:

My residence, post office address and citizenship are stated below next to my name:

I believe I am the original, first, and sole inventor (if only one name is listed below) or an original, first, and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled:

TITLE OF INVENTION

**BIOLOGICAL COMPOSITIONS, COMPONENTS THEREOF AND USES
THEREFOR**

the specification of which is attached hereto unless the following box is checked

was filed on November 10, 1999 as Application No. _____
or PCT Application No. PCT/AU99/00993 and amended on _____

(if applicable).

I hereby state that I have reviewed and understand the contents of the above-identified specification, including the claims, as amended by any amendment referred to above.

I acknowledge the duty to disclose information which is material to the examination of this application in accordance with 37 CFR §1.56.

I hereby claim foreign priority benefits under 35 U.S.C. §119(a)-(d) or §365(b) of any foreign application(s) for patent or inventor's certificate, or §365(a) of any PCT international application which designated at least one country other than the United States, listed below and have also identified below any foreign application for patent or inventor's certificate or PCT International application having a filing date before that of the application on which priority is claimed:

PRIOR FOREIGN/PCT APPLICATION(S)

COUNTRY/OFFICE	APPLICATION NO.	DATE OF FILING	PRIORITY CLAIMED
AU	PP 7060	November 11, 1998	<input checked="" type="checkbox"/> YES <input type="checkbox"/> NO

I hereby claim the benefit under 35 U.S.C. §119(e) of any United States provisional application(s) listed below.

PROVISIONAL APPLICATION NUMBER

DATE OF FILING

I hereby claim the benefit under 35 U.S.C. §120 of any United States application(s) or §365(c) of any PCT International application(s) designating the United States, listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States application in the manner provided by the first paragraph of 35 U.S.C. §112, I acknowledge the duty to disclose material information as defined in 37 CFR §1.56 which became available between the filing date of the prior application and the national or PCT international filing date of this application:

**PRIOR U.S. APPLICATIONS OR PCT INTERNATIONAL APPLICATIONS
DESIGNATING THE U.S. FOR BENEFIT UNDER 25 U.S.C. §120**

Status (check one)

Application Serial No.	Date of Filing	Patented	Pending	Abandoned
		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

5

And I hereby appoint Arthur H. Seidel, Registration No. 15,979; Gregory J. Lavorgna, Registration No. 30,469; Daniel A. Monaco, Registration No. 30,480; Thomas J. Durling, Registration No. 31,349; and John J. Marshall, Registration No. 29,671, my attorneys or agents with full power of substitution and revocation, to prosecute this application and to transact all business in the Patent and Trademark Office connected therewith.

Address all correspondence to Drinker Biddle & Reath LLP, One Logan Square, 18th & Cherry Streets, Philadelphia, PA 19103-6996. Address all telephone calls to Daniel A. Monaco, (215) 988-3312 (telefax: (215) 988-2757).

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code, and that such willful false statements may jeopardize the validity of the application or any patent issuing thereon.

FULL NAME OF SOLE OR FIRST INVENTOR

HOD

STEPHEN

(GIVEN NAME)

ALISTER

(MIDDLE INITIAL OR NAME)

LOCARNINI

(FAMILY OR LAST NAME)

Inventor's signature: 

Date: 1st June 2001

Country of Citizenship: Australia AUX

Residence: Victoria Australia

(City)

(State or Foreign Country)

Post Office Address: 13 Carlisle Avenue
East St. Kilda
Victoria 3183 Australia

FULL NAME OF SOLE OR SECOND INVENTOR

JOSEPH

(GIVEN NAME)

TORRESI

(FAMILY OR LAST NAME)

Inventor's signature: 

Date: 

Country of Citizenship: Australia

Residence: Victoria Australia

(City)

(State or Foreign Country)

Post Office Address: 9 Barriedale Court
Etham
Victoria 3095 Australia

FULL NAME OF SOLE OR FIRST INVENTOR

STEPHEN

(GIVEN NAME)

ALISTER

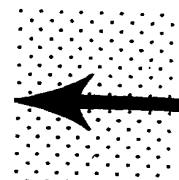
(MIDDLE INITIAL OR NAME)

LOCARNINI

(FAMILY OR LAST NAME)

Inventor's signature: X 

Date: X 1st June 2001



Country of Citizenship: Australia

Residence: Victoria

(City)

Australia

Post Office Address:

13 Carlisle Avenue

(State or Foreign Country)

East St. Kilda

Victoria 3183 Australia

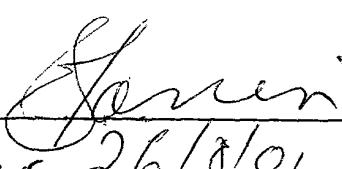
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JOSEPH

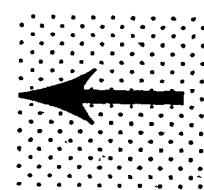
(GIVEN NAME)

TORRESI

(FAMILY OR LAST NAME)

Inventor's signature: X 

Date: X 26/8/01



Country of Citizenship: Australia

ALEX

Residence: Victoria

Australia

(City)

(State or Foreign Country)

Post Office Address:

9 Barriedale Court

Etham

Victoria 3095 Australia

FULL NAME OF SOLE OR THIRD INVENTOR

3-00 LINDA

(GIVEN NAME)

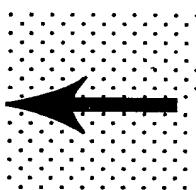
(MIDDLE INITIAL OR NAME)

EARNEST-SILVEIRA

(FAMILY OR LAST NAME)

Inventor's signature: Earnest

Date: 6/6/01



Country of Citizenship: Singapore SGX

Residence: Victoria Australia

(City)

(State or Foreign Country)

Post Office Address: 17 Kimberely Way
Bulleen
Victoria 3105 Australia

FULL NAME OF SOLE OR FOURTH INVENTOR

ANGELINE

(GIVEN NAME)

INGRID

(MIDDLE INITIAL OR NAME)

BARTHOLOMEUSZ

(FAMILY OR LAST NAME)

Inventor's signature: A. Bartholomeusz

Date: 31/05/01

Country of Citizenship: Australia

Residence: Victoria Australia

(City)

(State or Foreign Country)

Post Office Address: 64 Miller Street
Carnegie
Victoria 3163 Australia

FULL NAME OF SOLE OR THIRD INVENTOR

LINDA

(GIVEN NAME)

EARNEST-SILVEIRA

(FAMILY OR LAST NAME)

Inventor's signature: _____

Date: _____

Country of Citizenship: Singapore

Residence: Victoria Australia

(City)

(State or Foreign Country)

Post Office Address: 17 Kimberely Way
Bulleen
Victoria 3105 Australia

FULL NAME OF SOLE OR FOURTH INVENTOR

ANGELINE

(GIVEN NAME)

INGRID

(MIDDLE INITIAL OR NAME)

BARTHOLOMEUSZ

(FAMILY OR LAST NAME)

Inventor's signature: A. Bartholomeusz

Date: 31/05/01

Country of Citizenship: Australia AUSTRALIA

Residence: Victoria Australia

(City)

(State or Foreign Country)

Post Office Address: 64 Miller Street
Carnegie
Victoria 3163 Australia